Differential Patterns of Glucose-Induced Electrical Activity and Intracellular Calcium Responses in Single Mouse and Rat Pancreatic Islets

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Although isolated rat islets are widely used to study in vitro insulin secretion and the underlying metabolic and ionic processes, knowledge on the properties of glucose-induced electrical activity (GIEA), a key step in glucose-response coupling, has been gathered almost exclusively from microdissected mouse islets. Using a modified intracellular recording technique, we have now compared the patterns of GIEA in collagenase-isolated rat and mouse islets. Resting membrane potentials of rat and mouse β-cells were approximately −50 and −60 mV, respectively. Both rat and mouse β-cells displayed prompt membrane depolarizations in response to glucose. However, whereas the latter exhibited a bursting pattern consisting of alternating hyperpolarized and depolarized active phases, rat β-cells fired action potentials from a nonoscillating membrane potential at all glucose concentrations (8.4–22.0 mmol/l). This was mirrored by changes in the intracellular Ca²⁺ concentration ([Ca²⁺]), which was oscillatory in mouse and nonoscillatory in rat islets. Stimulated rat β-cells were strongly hyperpolarized by diazoxide, an activator of ATP-dependent K⁺ channels. Glucose evoked dose-dependent depolarizations and [Ca²⁺]i increases in both rat (EC₅₀ 5.9–6.9 mmol/l) and mouse islets (EC₅₀ 8.3–9.5 mmol/l), although it did not affect the burst plateau potential in the latter case. We conclude that there are important differences between β-cells from both species with respect to early steps in the stimulus-secretion coupling cascade based on the following findings: 1) mouse β-cells have a larger resting K⁺ conductance in 2 mmol/l glucose, 2) rat β-cells lack the compensatory mechanism responsible for generating membrane potential oscillations and holding the depolarized plateau potential in mouse β-cells, and 3) the electrical and [Ca²⁺]i dose-response curves in rat β-cells are shifted toward lower glucose concentrations. Exploring the molecular basis of these differences may clarify several a priori assumptions on the electrophysiological properties of rat β-cells, which could foster the development of new working models of pancreatic β-cell function. Diabetes 49:2028–2038, 2000

Isolated rat pancreas and islet preparations have been widely used to study in vitro insulin secretion and the underlying metabolic and ionic processes. One of the reasons is that, among small laboratory animals, rats exhibit a biphasic pattern of glucose-induced insulin release very similar to that of humans (1). Yet, another important reason is that the rat pancreas is an abundant source of isolated islets (hundreds of islets can be obtained from a single animal by successful enzymatic digestion), an essential condition for the usefulness of many of the techniques employed in islet studies.

Generation of electrical activity is a key step in glucose-induced insulin release, and it is widely thought to represent the primary mechanism by which Ca²⁺ is imported into the β-cell cytosol, causing an increase in the intracellular Ca²⁺ concentration ([Ca²⁺]) and triggering several processes that ultimately result in insulin exocytosis (2). Microelectrode recording of membrane potential from rat islets has been hampered by extreme technical difficulties related to the fact that these islets are small, hard to visualize under the dissecting microscope, and, thus, virtually impossible to isolate in a form amenable to fixation in a conventional electrophysiological chamber. Because these methodological constraints do not apply to mouse islets, intracellular recording of membrane potential has been traditionally carried out using microdissected mouse islets. Islet researchers have therefore been compelled to rely on the latter recordings for studies involving rat preparations. The validity of this extrapolation can, however, be brought into question by growing evidence that both confirmed the original observation (3) that rat and mouse islets exhibit differences in the pattern of glucose-induced insulin release and suggested the existence of clear differences in the pathways underlying stimulus-response coupling. For example, rat islets exhibit an enhanced production of (and sensitivity to) intracellular cAMP when stimulated with glucose (4). Also, glucose stimulation of rat islets induces a five- to sixfold increase in phosphoinositide hydrolysis, in sharp contrast to mouse islets in which virtually no increase was observed (5). Moreover, in rat β-cells, short-term exposure to high glucose induces a potentiation (priming) of the response to subsequent stimulation (6); this phenomenon is absent in mouse β-cells (7).

In a rather preliminary report (8), β-cells from collagenase-isolated cultured rat islets have been claimed to display 7-mV depolarizations in response to 11 mmol/l glucose and concomitant membrane potential fluctuations lasting ~1 s. At this sugar concentration, β-cells from microdissected mouse

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Received for publication 22 February 2000 and accepted in revised form 14 August 2000.

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BSA, bovine serum albumin; [Ca²⁺]i, intracellular Ca²⁺ concentration; GIEA, glucose-induced electrical activity; K ATP, ATP-dependent K⁺ channel.
islets exhibit slow variations in membrane potential under
lying bursts of action potentials, with typical durations and fre-
quencies of several seconds and 1–6 min⁻¹, respectively (9).
We have now made a systematic effort to not only charac-
terize glucose-induced electrical activity in collagenase-isola-
ated rat islets, but to actually compare its pattern with that
determined from parallel experiments on isolated mouse
islets carried out under identical experimental conditions.
This was achieved by further developing the recording tech-
nique, as described by Ikeuchi et al. (8), that was based on
the use of suction glass pipettes to hold in place collagenase-isola-
ated islets. Because previous studies had demonstrated a
tight correlation between oscillatory electrical activity and
[Ca²⁺], changes (10), we have also compared rat and mouse
islets from the standpoint of its cytosolic Ca²⁺ responses to
glucose stimulation.

RESEARCH DESIGN AND METHODS

Islet preparation and culture. Female albino mice (Charles River) 3–to
6-months-old and weighing 30–40 g and 9- to 13-week-old female Wistar rats
weighing 200–300 g were killed by a blow to the head, followed by cervical dis-
location. The retinal, heart, and islet cells were isolated by collage-
ase digestion of the pancreas, in accord with established
procedures (11,12). Isolated islets were visualized under a stereo microscope
and selected by its characteristic round shape, smooth edges, and white color.
In some experiments, islets (henceforth designated as freshly isolated
islets) were used shortly (i.e., a few hours) after isolation. These islets were
typically incubated in either 11 mmol/l glucose (mouse) or 8.4 mmol/l glucose
(rat) for 2 h and were further incubated for at least 50 min in 2 mmol/l glucose
before the experiments. The physiological salt solution had the following
composition (mmol/l): 120 NaCl, 5 KCl, 25 NaHCO₃, 2.56 CaCl₂,1.13 MgCl₂, and
3% bovine serum albumin (BSA) (pH 7.4 after equilibration with 5% CO₂/95%
O₂ in an humidified air atmosphere at 37°C). In other experiments, islets
(henceforth designated as cultured islets) were kept in culture for 15–24 h in
an RPMI-1640-based medium supplemented with 10% fetal calf serum, anti-
obiotics (100 U/ml penicillin and 100 µg/ml streptomycin), and glucose (11 and
8.4 mmol/l for mouse and rat islets, respectively), pH 7.4. Other tissue culture
details have been described by Salgado et al. (12). Like the freshly isolated
islets, these islets were further incubated for at least 50 min in 2 mmol/l glu-
cose before the experiments.

Electrophysiological experiments. The membrane potential was recorded
from collagenase-isolated islets using a high-impedance amplifier as previously
reported (13). Islets were then transferred to a fast perfusion chamber (vol
40 µl), in which they were held in place with the help of a suction pipette.
This procedure consisted of a fire-polished glass pipette filled with a physiologi-
ical salt solution and joined to a fine plastic tubing to which a gentle negative
pressure was applied with the help of a syringe. The suction pipette was posi-
tioned with a micromanipulator. The perfusion solution was the same as the
above physiological salt solution, except that it contained no BSA, was sup-
plemented with different concentrations of glucose as required, and was con-
continuously gassed with 95% O₂/5% CO₂ for a final pH of 7.4. This solution was deliv-
ered to the chamber at ~1.5 ml/min (37°C). Solution changes were made with
the help of a stop-cock valve. The time required for complete solution
exchange at the chamber level was ~8 s (not corrected for in the figures). Islet
β-cells were impaled with high resistance (100–200 MΩ) glass microelec-
trodes. Input resistance to the β-cell membrane was assessed by injecting rec-
tangular pulses of current (0.1 nA) through the recording microelectrode
after null-bridge offset compensation in extracellular solution of the voltage
deflections due to microelectrode resistance, as reported (14). Other electro-
physiological details have been described by Rosário et al. (13). Impalement
of β-cells was identified by the characteristic cell depolarization in response
to 11 mmol/l glucose or tolbutamide.

Microfluorometry. The [Ca²⁺], was measured from single islets using the flu-
orescent Ca²⁺ indicator fura-2 as described previously (11,12). Briefly, islets were
incubated with a physiological salt solution supplemented with 3% BSA, 2
mM CaCl₂, 11 mM glucose, and 5 µmol/l fura-2 AM for 15–30 min at 37°C. The loaded
islets were then transferred to a perfusion chamber (30 µl vol, 1.5 ml/min flow rate),
placed on the stage of an inverted fluorescence microscope, and allowed to
attach to a poly-L-lysine–coated coverslip (used as a chamber bottom) for 10–15
min before starting the experiments. Solution changes were made with a
stop-cock, and there was a delay to the chamber of 11 s (not compensated for
in the figures). The [Ca²⁺], was measured using a dual excitation microflu-
rescence system (Deltascan; Photon Technology International, Princeton,
NJ). Fura-2 was excited at 340/380 nm, and the fluorescence was detected using
an interference filter centered at 510 nm. The data were corrected for back-
ground fluorescence and were acquired at 10 Hz by a computer. The fluores-
cence ratio F₅₁₀/F₃₈₀ was converted into [Ca²⁺], using the calibration
equation of Gryniewicz et al. (15), as described previously (11).

RESULTS

Figure 1 depicts representative examples of electrophysio-
logical experiments in which freshly isolated rat and mouse
islets (islets 1 and 2 in Fig. 1A and islet 1 in Fig. 1B) were ini-
tially exposed to a subthreshold glucose concentration
(2 mmol/l) and subsequently challenged with 11 mmol/l glu-
cose. We have also recorded membrane potential from islets
(e.g., islet 3 in Fig.1A and islet 2 in Fig.1B) that have been
maintained in culture for 18–24 h to allow full recovery from
the isolation procedure.

Resting membrane potential and [Ca²⁺]. Rat islet β-cells
exhibited a more positive resting membrane potential than
mouse islets. Indeed, as shown in Fig. 1C, the average mem-
brane potential recorded from freshly isolated rat and mouse
islets in 2 mmol/l glucose was ~50 ± 2 and ~60 ± 3 mV, respec-
tively (n = 8–13 islets, P < 0.05). This difference is unlikely to
be a consequence of the isolation procedure, because cultured
rat islets were also more depolarized than cultured mouse
islets in 2 mmol/l glucose (average resting membrane potential
~52 ± 3 and ~59 ± 2 mV, respectively; n = 8–10 islets; P < 0.05). The fact that rat
β-cells are relatively more depolarized is consistent with the display of higher levels of noise (flick-
ering potential) by membrane potential recordings in these
cells compared with mouse β-cells (Fig. 1). At this point, it
should be noted that exposing rat β-cells to diazoxide, an activ-
ator of ATP-dependent K⁺ (KATP) channels (16), brought the
membrane potential to levels significantly more negative
(approximately ~20 mV) than those recorded in 2 mmol/l glucose
alone (Fig. 2A).

We have recorded the [Ca²⁺], from whole collagenase-isola-
ted islets using the fluorescent Ca²⁺ indicator fura-2 (Fig. 4).
As shown in Fig. 4C, there was no significant difference be-
tween resting [Ca²⁺], in rat and mouse islets (67 ± 4 and 55 ±
8 nmol/l, respectively, for freshly isolated islets, n = 8–18;
61 ± 4 and 62 ± 6 nmol/l, respectively, for cultured islets, n = 7–18;
P > 0.05 in both cases).

Effects of 11 mmol/l glucose on membrane potential and
[Ca²⁺]. Fig. 1B shows that glucose (11 mmol/l) depolarized the
β-cell membrane and triggered a bursting electrical activity
(slow membrane potential waves with action potentials firing
from a plateau potential of approximately ~35 mV) in col-
lagenase-isolated mouse islets. This phenomenon resembles
the typical bursting electrical activity recorded from
microdissected mouse islets (17).

Raising the glucose concentration from 2 to 11 mmol/l
evoked a prompt membrane depolarization in all rat islets
tested (Fig. 1A). This finding, however, differed markedly
from the glucose response displayed by mouse islets, since
there were no oscillations in membrane potential associ-
ated with bursting electrical activity. The amplitude of the glu-
cose-induced depolarization was variable from islet to islet
and, for the most part, appeared to increase along the first
few minutes of stimulation. For freshly isolated islets, the
FIG. 1A and B. Glucose-induced electrical activity in rat and mouse β-cells. Glucose concentration in the perfusion medium was raised from 2 (G2) to 11 mmol/l (G11) as indicated by the horizontal bars. A: Typical intracellular recordings from freshly isolated (islets 1 and 2) and cultured (islet 3) rat islets. B: Typical intracellular recordings from freshly isolated (islet 1) and cultured (islet 2) mouse islets. Details of electrical activity are shown beneath the main records on an expanded time scale. Islets were isolated by collagenase digestion in all cases. The membrane potential of stimulated β-cells was measured at the foot of the spikes.
maximal amplitude of depolarization, defined as the difference between membrane potential at the steady state in 11 mmol/l glucose (foot of the spikes) and resting membrane potential in 2 mmol/l glucose, averaged 23 ± 5 mV (n = 6 islets). This is of the same order of magnitude as the maximal amplitude of depolarization reached in mouse islets (difference between plateau potential and resting membrane potential in 2 mmol/l glucose: 18 ± 2 mV, n = 6 islets) (Fig. 1C).

Membrane depolarization was accompanied by the appearance of action potentials in all rat islets examined. However, the amplitude of these spikes was clearly variable from islet to islet (and often within a particular islet), depending on the membrane potential at the foot of the spikes. This is especially evident in islet 1 (Fig. 1A, expanded trace a), in which well-defined voltage fluctuations (i.e., clearly in excess of basal noise levels) could be observed only over a relatively narrow membrane potential range (approximately –40 to –25 mV). For other islets, the depolarizing effect of glucose was slower and less intense (membrane potentials at the foot of the spikes were more negative than –30 mV at all times), and vigorous spiking persisted throughout an extended interval, albeit with a tendency to decrease in amplitude as the membrane depolarized (e.g., islet 2 in Fig. 1A). Figure 3A (left histogram) relates spike amplitude to membrane potential at the foot of the spikes (data collected from five experiments in which rat islets were stimulated with 11 mmol/l glucose). This analysis shows that spike amplitude increased with membrane potential in the range –50 to –35 mV; spike amplitude decreased for larger depolarizations and became close to basal noise levels for membrane potential values more positive than –25 mV. For mouse islets, the average spike amplitude in 11 mmol/l glucose showed a tendency to decrease with membrane potential in the range

![Image](image_url)
–42.5 to –30 mV and was of the same order of magnitude as that measured from rat islets (Fig. 3A, right histogram).

We have calculated maximum depolarization and repolarization rates of rat and mouse β-cell action potentials by determining the time derivatives of the membrane potential from expanded traces in 11 mmol/l glucose (range of membrane potential values at the foot of the spikes –35 to –33 mV). Figure 3B shows that maximum spike depolarization and repolarization rates were smaller for rat β-cells (average maximum depolarization rates pooled from three similar experiments on rat and mouse islets 0.35 ± 0.02 and 0.67 ± 0.05 V/s, respectively; average maximum repolarization rates –0.24 ± 0.01 and –0.49 ± 0.04 V/s, respectively). The spike depolarization rate in rat β-cells is much lower than would be expected for Na+ action potentials (or action potentials with a mixed contribution of Na+ and Ca2+ channels) (18), suggesting that it is mainly mediated by the activation of voltage-sensitive Ca2+ channels. Accordingly, Fig. 3C shows that the L-type Ca2+ channel blocker nifedipine abolished the rat β-cell electrical activity induced by 5.6 mmol/l glucose.

We have investigated the effect of 11 mmol/l glucose on [Ca2+]i, as determined from whole mouse and rat islets. Glucose induced a multiphasic [Ca2+]i response from mouse islets (Fig. 4B): the [Ca2+]i transiently decreased to levels below baseline, increased sharply to a peak, and oscillated from an elevated plateau for the remainder of the stimulation. These [Ca2+]i oscillations originate from oscillatory Ca2+ influx associated with bursting electrical activity (10). As shown in Fig. 4A, the initial effect of glucose on rat islets was to lower the [Ca2+]i; subsequently, the [Ca2+]i increased to a peak and either maintained elevation at near peak levels or decreased slowly to a plateau. High-frequency [Ca2+]i fluctu-
Associations, reminiscent of the spiking activity displayed by some rat islets, were sporadically recorded at the beginning of the \([\text{Ca}^{2+}]_i\) rise (Fig. 4A, islet 1; representative of 7 of 22 experiments). Interestingly, peak \([\text{Ca}^{2+}]_i\) values were reached between 4 and 6 min of stimulation, regardless of the animal species (286 ± 17 and 338 ± 22 s for mouse and rat islets, respectively; \(n = 11–22\) islets). This matches the time course of the first phase of glucose-induced insulin release (19).

There were some differences between freshly isolated and cultured mouse islets with respect to their responsiveness to 11 mmol/l glucose. For example, while cultured islets displayed a typical biphasic response consisting of continuous spiking followed by regular bursting (Fig. 1B, islet 2; representative of six islets), most of the freshly isolated islets examined (three of four) exhibited instead an initial pattern of higher frequency bursting (Fig. 1B, islet 1). It is also noteworthy that the amplitude of

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**FIG. 4.** Effect of 11 mmol/l glucose on \([\text{Ca}^{2+}]_i\), in rat and mouse islets. Glucose concentration was changed between 2 and 11 mmol/l (G11) as indicated by the arrows. The vertical scale represents the \([\text{Ca}^{2+}]_i\), as measured from whole islets by a dual excitation epifluorescence procedure (RESEARCH DESIGN AND METHODS). A: Typical \([\text{Ca}^{2+}]_i\) recordings from freshly isolated (islets 1 and 2) and cultured (islet 3) rat islets. B: Typical \([\text{Ca}^{2+}]_i\) recordings from freshly isolated (islet 1) and cultured (islet 2) mouse islets. C: Average \([\text{Ca}^{2+}]_i\) responses to glucose stimulation in rat and mouse islets. Data were pooled from the experiments depicted in A and B and similar experiments. Islets were either used fresh after collagenase isolation (Fresh) or kept in culture for 18–24 h (Culture), as explained in RESEARCH DESIGN AND METHODS. Measurements were made either in 2 mmol/l glucose or at the steady state in 11 mmol/l glucose, as indicated below the columns. The plotted values represent the average \([\text{Ca}^{2+}]_i\), at 15–18 min during stimulation. Data are presented as means ± SE (the number of experiments performed is given above each column in parentheses).
[Ca\(^{2+}\)]\(_i\) oscillations is higher in cultured compared to freshly isolated islets (Fig. 4B; islets 2 and 1, respectively). As a result, the average amplitude of the [Ca\(^{2+}\)]\(_i\) response, assessed by averaging out the [Ca\(^{2+}\)]\(_i\) at the steady state, was notably higher for cultured islets (Fig. 4C). However, as shown in Fig. 1C, maximal amplitude of depolarization reached in cultured islets did not differ from that in freshly isolated islets.

Freshly isolated and cultured rat islets displayed a similar responsiveness to 11 mmol/l glucose, as shown in Figs. 1C and 4C. For example, maximal amplitude of depolarization recorded from cultured islets averaged 22 ± 6 mV (n = 4 islets; not significantly different from freshly isolated islets [P > 0.05]). The amplitude of the [Ca\(^{2+}\)]\(_i\) response in freshly isolated rat islets (55 ± 3 nmol/l, n = 18 islets) was also similar to that recorded from cultured islets (62 ± 4 nmol/l, n = 18 islets) (as shown in Fig. 4A, in which islets 1 and 2 and islet 3 are examples of freshly isolated and cultured islets, respectively).

**Dose-dependent effect of glucose on membrane potential and [Ca\(^{2+}\)]\(_i\).** Rat islets have a lower threshold for glucose-induced insulin release (20, 21). It is therefore conceivable that both the membrane potential and the [Ca\(^{2+}\)]\(_i\) responses might be near maximal at 11 mmol/l glucose, thus accounting for the lack of sustained oscillations at the steady state for this glucose concentration. We have assessed this possibility by continuously recording the membrane potential and [Ca\(^{2+}\)]\(_i\) in response to stepwise increases in glucose concentration.

Figure 5A depicts a typical membrane potential recording from a rat islet stimulated with increasing glucose concentrations in the range 2–16.7 mmol/l. Raising the sugar concentration from 2 to 5.6 mmol/l evoked a multiphasic response, characterized by an early membrane depolarization and concomitant surge of spikes, followed by a slight hyperpolarization without spiking activity; a few minutes later, the cell depolarized to a level 10 mV above resting membrane potential, and spiking activity resumed. Further raising the glucose concentration to 8.4, 11, and 16.7 mmol/l evoked progressive and slow membrane depolarizations, bringing membrane potential at the foot of the spikes to different levels in the range –40 to –25 mV, as a consequence, spike amplitude decreased progressively and became residual at the highest glucose concentration tested. An increase in spike frequency
is also apparent at each step, especially for intermediate glucose concentrations (e.g., 0.9, 1.2, and 1.6 s⁻¹ for 5.6, 8.4, and 11 mmol/l, respectively, in the experiment depicted in Fig. 5A). The membrane potential at the foot of the spikes, measured at the steady state for each glucose concentration, was pooled across several experiments and plotted in Fig. 5C. This analysis yielded an average EC₅₀ for glucose-induced depolarization of the β-cell membrane of 5.9 mmol/l (average membrane potential at saturating glucose −33 ± 3 mV, n = 7 islets).

The effect of glucose on membrane potential of rat β-cells was matched by appreciable [Ca²⁺]ᵢ increases at all glucose concentrations, starting with 5.6 mmol/l (Fig. 5B). The average [Ca²⁺]ᵢ, at the steady state for each glucose concentration was measured from different islets and plotted in Fig. 5C, yielding an EC₅₀ of 6.9 mmol/l.

Figure 6A depicts the typical effect of raising glucose concentration on membrane potential of β-cells from mouse islets. In sharp contrast to rat, mouse β-cells exhibited a pattern of bursting electrical activity at all glucose concentrations in the range 8.4–22 mmol/l. Similar to microdissected mouse islets (22), raising the sugar concentration in this range evoked a dose-dependent increase in fractional active-phase duration (Fig. 6C). As a consequence, there was a dose-dependent increase in average spike frequency. The EC₅₀ values, calculated from these plots, were 9.5 and 8.3 mmol/l, respectively. Importantly, there were no consistent drifts in either plateau or silent-phase potential as the glucose concentration was raised (average plateau potential in 8.4, 16.7, and 22 mmol/l glucose –38 ± 1, –38 ± 1, and –38 ± 1 mV, respectively; n = 3 islets). It is also worth noting that 5.6 mmol/l glucose depolarized the β-cell membrane by 10 mV without evoking sustained electrical activity.

Fig. 6B shows that, in mouse islets, glucose evoked [Ca²⁺]ᵢ oscillations throughout the intermediate-to-high concentration range (11–22 mmol/l). Moreover, glucose increased the duration of these oscillations in a dose-dependent fashion (3.7 ± 0.1, 9 ± 1, and 11 ± 1 s at 11, 16.7, and 22 mmol/l glucose, respectively; n = 11–23 oscillations for each concentration). There was, however, no consistent change in either peak [Ca²⁺]ᵢ or the [Ca²⁺]ᵢ at the trough of the oscillations. We have estimated mean [Ca²⁺]ᵢ by averaging out the [Ca²⁺]ᵢ through-
out the last 120-s period at each glucose concentration. The EC50 value, estimated from a plot of average [Ca2+]i versus glucose concentration, was 8.7 mmol/l (Fig. 6C).

**Effect of glucose on β-cell input resistance.** We injected rectangular pulses of hyperpolarizing current (0.1 nA) to assess the effect of glucose stimulation on input resistance to the rat β-cell membrane (Fig. 2B). Input resistance was ~108 MΩ at 2 mmol/l glucose, and it increased by 90 and 190% when the cells were stimulated with 5.6 and 16.7 mmol/l glucose, respectively (Fig. 2C). These values compare well with the respective values for mouse β-cells (23).

**DISCUSSION**

Using an improved intracellular recording technique, we have shown that β-cells from rat islets of Langerhans have stable negative membrane potential levels at stimulatory glucose concentrations and that they become depolarized for glucose loads higher than 5.6 mmol/l. Furthermore, glucose-induced depolarization is accompanied by a pronounced rise in cell-input resistance and is antagonized by the KATP channel activator diazoxide. This is consistent with KATP channels being inhibited by glucose metabolism, an essential characteristic of the glucose-response-coupling mechanism that has been proposed for pancreatic β-cells of different animal species (24). We have also shown that membrane depolarization in rat β-cells is often accompanied by the firing of action potentials, albeit with an intensity and pattern that are clearly dependent on the extent of depolarization. Specifically, spiking activity is often negligible for membrane potential values more negative than ~40 mV. This is consistent with membrane depolarization activating high-threshold voltage-sensitive Ca2+ channels, most likely of the L-type, because glucose-induced electrical activity is impaired by the dihydropyridine nifedipine. Massive Ca2+ influx associated with the activation of these channels likely accounts for the pronounced [Ca2+]i changes displayed by rat islets.

The pattern of glucose-induced electrical activity recorded from rat islet β-cells differs from that of mouse β-cells in a number of important points. Whereas the latter exhibit a typical bursting pattern over an extended glucose concentration range (8.4–22.0 mmol/l) (as found in this work and in the study by Atwater et al. [23]), rat β-cells fire action potentials from a nonscillating membrane potential, regardless of the sugar concentration. This is also evident from the standpoint of [Ca2+]i, which is oscillatory in mouse and nonscillatory in rat islets. Therefore, it should be emphasized that, in mouse β-cells, glucose increases the duration of the slow membrane potential waves underlying bursting electrical activity without affecting plateau potential and, hence, the characteristics of the action potentials (23); as a result, there is a dose-dependent increase in the duration (not the amplitude) of the associated [Ca2+]i oscillations. In rat β-cells, in contrast, glucose depolarizes the membrane at the foot of the spikes in a dose-dependent way and more than likely increases the opening probability of the noninactivating fraction of voltage-sensitive Ca2+ channels. This action probably explains why the signaling mechanism in rat islets involves changes in the amplitude of the [Ca2+]i signals.

Different groups have reported varying patterns of [Ca2+]i and insulin responses of rat islets to glucose stimulation. Using single collagenase-isolated islets, Martin et al. (25) reported the absence of a [Ca2+]i oscillatory pattern in 95% of the islets stimulated with 11–16.7 mmol/l glucose, whereas Longo et al. (26) and Bergsten et al. (27,28) presented evidence for sustained [Ca2+]i, and insulin oscillations, respectively, in the presence of 10–11 mmol/l glucose. In vitro multi-islet preparations have also been shown to undergo sustained insulin pulsatility in response to 5.5–16.7 mmol/l glucose, albeit with a marked reduction in frequency compared with pulsatile insulin release from single islets (26,29,30). The reason for these apparent discrepancies is not known but could be related to specificities in islet handling (e.g., fresh versus cultured islets, glucose concentration in the culture/preincubation medium and composition of the perfusion solutions). However, it is worth noting that we did not find major differences between fresh and cultured islets in this study. The situation in mouse islets appears to resemble that of human and pig islets, which have been consistently shown to oscillate in response to 11–16.7 mmol/l glucose (31–33). Other parameters of β-cell function have been reported to exhibit marked interspecies differences. For instance, dog and rodent β-cells fire Na+- and Ca2+-dependent action potentials, respectively, whereas human β-cells display patterns of electrical activity with varying contributions from voltage-sensitive Na+ and Ca2+ channels (18,34,35).

Recent patch-clamp studies of β-cells in intact mouse islets (36) have revealed the presence of a small voltage-activated K+ conductance, reminiscent of the glucose-activated outward K+ current described by Rojas et al. (37). This current is strictly dependent on stimulation of Ca2+ influx, develops slowly in response to depolarizing pulses, and has the required size to counteract the depolarizing effect of KATP channel inhibition by glucose. It may therefore represent the primary mechanism by which the burst plateau potential is set at a rather constant level (approximately ~35 mV), irrespective of the sugar concentration. In isolated rat β-cells, in turn, glucose failed to stimulate net outward current (37), suggesting that either the cells lack the putative Ca2+-activated K+ channels or that the size of the Ca2+ currents is insufficient to activate these channels to the extent required to regulate the membrane potential. Thus, lack of this compensatory mechanism in rat islet β-cells may explain the fact that glucose depolarizes the cells in a dose-dependent fashion. It should be noted, however, that we cannot rule out the possibility that alternative ionic mechanisms might play a role in this depolarization. For example, Na+/Ca2+ exchange activity has been recently reported to be ~50% higher in rat β-cells (38,39). When operating in the forward mode, this system exchanges three Na+ ions for one Ca2+, thus providing the net transfer to the cytoplasm of one positive charge per cycle (40). This would cause rat β-cells to be somewhat more depolarized than mouse β-cells, in agreement with the results from our study.

There is no appreciable difference between mouse and rat β-cells with respect to the amplitude of the action potentials, provided that care is taken to match the membrane potential at the foot of the spikes. There is, however, a marked difference concerning the respective rates of depolarization and repolarization: spikes are clearly slower in rat, as evidenced by comparing its first derivatives. It has been proposed that the maximal inward and outward currents can be roughly estimated from the maximal derivatives of the ascending and descending phases of the action potentials, respectively (23).

Therefore, we hypothesize that the size of the Ca2+ currents responsible for the depolarizing phase of the action potentials...
is generally diminished in rat compared with mouse β-cells. Reduced Ca\textsuperscript{2+} loads arising from the activation of voltage-sensitive Ca\textsuperscript{2+} channels may in turn lower the repolarization rate of the action potentials in rat β-cells, assuming that this is partially controlled by Ca\textsuperscript{2+}-activated K\textsuperscript{+} channels.

Yet another difference is that average resting membrane potential in rat islet β-cells (2 mmol/l glucose) is 7–10 mV more positive than in mouse β-cells. Two hypotheses may be put forward to explain this difference: mouse β-cells have a larger resting K\textsuperscript{+} conductance (higher density of K\textsubscript{ATP} channels or reduced fraction of K\textsubscript{ATP} channels inhibited by 2 mmol/l glucose); alternatively, rat β-cells may have a larger resting Na\textsuperscript{+} conductance. In a recent patch-clamp study (performed whole-cell recording configuration, 10 mV pulses), Gopel et al. (41) reported K\textsubscript{ATP} channel currents of ~26 pA for mouse β-cells in the absence of glucose. Taking into account cell capacitance, this corresponds to a K\textsubscript{ATP} channel density of ~4 pA/pF. In a similar study conducted using rat β-cells, Hughes et al. (42) reported K\textsubscript{ATP} channel densities of 4.6 pA/pF. This suggests that K\textsubscript{ATP} channel density is not significantly different between the two species and is therefore in agreement with our finding that diazoxide brings the resting membrane potential of rat β-cells to a level close to that of mouse β-cells.

We have recalculated available patch-clamp data (43) to assess the dependence of K\textsubscript{ATP} channel activity on free ATP levels in excised inside-out patches of mouse β-cell membranes. This analysis yielded an IC\textsubscript{50} for K\textsubscript{ATP} channel inhibition of 1.4 mmol/l, a value close to that determined by Ashcroft and Kakei (44) for rat β-cells (2 mmol/l). It is likely, then, that the difference between mouse and rat β-cells might be explained by different cytosolic ATP/ADP ratios rather than by a different sensitivity to intracellular ATP. Available measurements of cytosolic intracellular purine nucleotides do not accurately reflect the cytosolic pool, implying that cytosolic ATP/ADP ratios cannot be readily estimated in β-cells. However, it is worth noting that raising the glucose concentration from 0 to 3 mmol/l increased the ATP/ADP ratio (calculated from total ATP and ADP levels) by 10 and 70% in mouse and rat islets, respectively (45–48). This finding reinforces the view that in rat β-cells exposed to a subthreshold glucose concentration, a higher proportion of K\textsubscript{ATP} channels is inhibited by a higher ATP/ADP ratio.

We have determined the dose dependency of rat and mouse β-cell function, using a number of electrophysiological and [Ca\textsuperscript{2+}] parameters. The estimated EC\textsubscript{50} values for these parameters lay in the range 5.9–6.9 and 8.3–9.5 mmol/l, respectively, indicating that the dose-response curves for rat β-cells are shifted to lower glucose concentrations compared with mouse β-cells. This is in keeping with published data for glucose-induced insulin secretion (20,21) and may relate to the fact that low-K\textsubscript{o} hexokinase activity accounts for a higher fraction of total glucose phosphorylating activity in rat compared with mouse β-cells (49,50). The differential responses to the lower depolarizing glucose concentration used in our study (5.6 mmol/l) are particularly noteworthy: rat β-cells display a sustained spiking activity, whereas mouse β-cells do not. This is consistent with rat islets having a lower threshold for glucose-induced insulin release.

In summary, our study shows that there are important differences between β-cells from rat and mouse islets, the two most widely used preparations in β-cell research, with respect to early steps in the stimulus-secretion coupling cascade. In particular, rat β-cells lack the compensatory mechanism responsible for generating membrane potential oscillations and for holding the depolarized plateau potential in mouse β-cells. This implies that, contrary to current views, rat islets or β-cells cannot be assumed to bear the electrophysiological properties of mouse islets or β-cells. Realizing the differences between β-cells from both species may lead to the development of new working models of pancreatic β-cell function.

ACKNOWLEDGMENTS

C.M.A. and A.P.S. were supported by the Praxis XXI Program (Portugal). This study was supported in part by Fundação para a Ciência e Tecnologia, Praxis XXI, and the Islet Research European Network Concerted Action of the Commission of the European Communities. We gratefully acknowledge the School of Medicine at the University of Coimbra for providing the facilities, which were essential for conducting this work.

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