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Abstract

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Desensitization of Mitochondrial Ca\textsuperscript{2+} and Insulin Secretion Responses in the Beta Cell*

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The role of mitochondria in the desensitization of insulin secretion was investigated. In rat pancreatic beta cells, both insulin secretion and mitochondrial [Ca\textsuperscript{2+}] increases were desensitized following two challenges with the mitochondrial substrate methyl succinate. In the beta cell line INS-1, similar results were observed when a 5-min interval separated two 5-min pulses. In contrast, ATP generation monitored in luciferase-expressing INS-1 cells was stimulated to the same extent during both exposures to methyl succinate. Succinate, like α-glycerophosphate, activates the electron transport chain at complex II. As a consequence, the mitochondrial membrane hyperpolarizes, promoting ATP synthesis and Ca\textsuperscript{2+} influx into the mitochondria through the uniporter. The mitochondrial desensitization was further studied in permeabilized INS-1 cells. Increasing extramitochondrial [Ca\textsuperscript{2+}] from 100 to 500 nM enhanced succinate oxidation 4-fold. At 500 nM Ca\textsuperscript{2+}, 1 mM succinate caused a blunted mitochondrial [Ca\textsuperscript{2+}] increase upon the second, compared with the first, stimulation. These effects were mimicked by α-glycerophosphate, and there was cross-desensitization between the two compounds. Succinate hyperpolarized the mitochondrial membrane during both the first and second applications. This suggests that the uniporter itself, rather than the respiratory chain, is desensitized. These results emphasize the key role of the mitochondria not only in the stimulation of insulin secretion, but also in its desensitization.

Desensitization is a common feature of cell biology in general and of insulin secretion in particular. However, the molecular mechanism of desensitization toward nutrient stimuli is poorly understood. Nesher and Cerasi (1) first observed that successive short stimuli with glucose or arginine in the isolated perfused rat pancreas resulted in the inhibition of the insulin secretory response to the second stimulus. Insensitivity of the pancreatic beta cell to glucose was reported in pancreata taken from hyperglycemic rats (2) and is found in several diabetic animal models (3). A reduced responsiveness of the pancreatic beta cell to glucose has also been described after prolonged exposure of beta cells to hexose in vitro (4, 5) or in human subjects (6). This desensitization phenomenon is distinguished from glucose toxicity, the latter being irreversible, whereas the former implies a reversible state of cellular refractoriness due to repeated exposures to an agonist (7). Desensitization can occur at any of the multiple steps coupling glucose recognition to insulin secretion, including the exocytotic process itself, as shown in permeabilized cells exposed to repeated Ca\textsuperscript{2+} pulses (8).

In the pancreatic beta cell, mitochondrial metabolism plays a pivotal role in the generation of signals coupling glucose recognition to insulin secretion (9–13). The main trigger of exocytosis is an increase in cytosolic Ca\textsuperscript{2+} concentration (for a review, see Ref. 12). In addition, Ca\textsuperscript{2+} controls several other cellular functions, among them mitochondrial metabolism (14–16). An increase in mitochondrial Ca\textsuperscript{2+} concentration ([Ca\textsuperscript{2+}]\textsubscript{m})\textsuperscript{,} following an elevation in cytosolic Ca\textsuperscript{2+} concentration ([Ca\textsuperscript{2+}]\textsubscript{c}), participates in the activation of the respiratory chain through stimulation of Ca\textsuperscript{2+}-sensitive NADH-generating dehydrogenases (15–20). NADH and FADH\textsubscript{2} transfer reducing equivalents to the respiratory chain, thereby ensuring adequate ATP synthesis (15). Transfer of reducing equivalents to the electron transport chain increases the mitochondrial membrane potential (ΔΨ\textsubscript{m}), which enhances the driving force for mitochondrial Ca\textsuperscript{2+} uptake mediated by a low affinity uniporter (21). This ΔΨ\textsubscript{m}-dependent Ca\textsuperscript{2+} entry permits an amplification of [Ca\textsuperscript{2+}]\textsubscript{m}, relative to [Ca\textsuperscript{2+}]\textsubscript{c}, further favoring the stimulation of the aforementioned dehydrogenases (22, 23). On the other hand, the hyperpolarization of the mitochondrial membrane exerts a negative feedback by lowering the oxygen consumption and the rate of H\textsuperscript{+} cycling (24, 25). In glucose-stimulated beta cells, insulin secretion is initiated by the activation of mitochondrial metabolism, leading to an increase in [Ca\textsuperscript{2+}], (10, 26, 27). Subsequently, the rise in [Ca\textsuperscript{2+}]\textsubscript{m} appears to be essential for the maintenance of metabolism-secretion coupling (12, 13). The partial reduction of glucose oxidation by blockade of the [Ca\textsuperscript{2+}]\textsubscript{c} increase (17, 28) may reflect a need for permissive [Ca\textsuperscript{2+}]\textsubscript{c} levels in optimal glucose-stimulated insulin secretion (29).

Using cells stably expressing the calcium-sensitive photoprotein aequorin targeted to the mitochondria, we have previously shown that desensitization of insulin secretion is associated with a parallel loss of the [Ca\textsuperscript{2+}]\textsubscript{m} response (23). These findings and other recent studies point to a pivotal role for the mitochondria in metabolism-secretion coupling (11, 17, 20, 30–32), not only as a relay in the metabolic cascade, but also as a primary source of an as yet unidentified factor triggering insulin exocytosis (13). The existence of this putative mitochondrial factor is further suggested by studies showing impaired glucose-stimulated insulin secretion in insulinoma cells depleted of the mitochondrial genome (33, 34).

To study the involvement of the mitochondria in the desen-

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1 The abbreviations used are: [Ca\textsuperscript{2+}]\textsubscript{m}, mitochondrial Ca\textsuperscript{2+} concentration; [Ca\textsuperscript{2+}]\textsubscript{c}, cytosolic Ca\textsuperscript{2+} concentration; ΔΨ\textsubscript{m}, mitochondrial membrane potential.
sition process, we have monitored insulin secretion and three parameters that reflect mitochondrial activation: 1) $\Delta \psi_m$, using the fluorescent probe rhodamine 123, 2) $[Ca^{2+}]_{im}$ employing a cell line stably expressing mitochondrial aequorin, and 3) generation of ATP using a cell line stably expressing cytosolic luciferase. The tricarboxylic acid cycle intermediate succinate was used as a mitochondrial substrate. Some of these experiments were performed in cells isolated from rat islets, whereas the remainder were in cells derived from the rat insulinoma cell line INS-1 (35). To study the dissociation between the $[Ca^{2+}]_{im}$ signal and ATP generation, experiments were performed in *Staphylococcus* α-toxin-permeabilized INS-1 cells, which permits the control of the mitochondrial environment with respect to cytosolic $[Ca^{2+}]$ and [ATP] (13). The results show that $[Ca^{2+}]_{im}$ increases and insulin secretion are strongly desensitized by mitochondrial substrates, whereas generation of ATP and $\Delta \psi_m$ activation are not. The study provides evidence that the mitochondrial Ca$^{2+}$ uniporter is desensitized, rather than the activation of the electron transport chain.

**EXPERIMENTAL PROCEDURES**

*Cell Culture—* INS-1 cells were cultured in RPMI 1640 medium as described previously (23, 35, 36). Stable clones of INS-1 cells expressing mitochondrial aequorin (22) were established (INS-1-LUC7) (36). Following two successive stable transfections, resistant clones were cultured with 250 $\mu$g/ml G418 and 100 $\mu$g/ml hygromycin B (Calbiochem). Pancreatic islet cells were isolated by collagenase digestion from male Wistar rats weighing ~200 g (17) and cultured free floating in RPMI 1640 medium for 2–4 days.

*Transient Transfection of Primary Cells—* Rat pancreatic islet cells were isolated as described above, trypsinized, and seeded on 13-mm diameter extracellular matrix-coated coverslips (Eldan, Jerusalem, Israel) at 4 x $10^5$ cells/ml in RPMI 1640 medium. Two days later, the cells were transfected with 10 $\mu$l of LipofectAMINE (Gibco BRL, Basel, Switzerland) and 1 $\mu$g of vector encoding mitochondrial targeted aequorin as described previously (17). Three days later, the cells were used for the experiments. This transfection resulted in 10–15% of cells being transfected as judged by immunofluorescence of the hemagglutinin tag incorporated at the N terminus of aequorin (22, 23).

*Permeabilization of Cells—* Attached INS-1 cells were grown on A431 extracellular matrix as described above. Cells were maintained in subconfluent monolayer for 3–4 days prior to transformation. The cells were then placed in a thermostatted microincubator (Medical Systems Corp., Greensville, VA) on an inverted microscope (Nikon Diaphot) with a 40× oil immersion objective. Fluorescence excitation was filtered at 485 nm, and emission was split at 505 nm and further filtered at 530 nm (Omega Optical Inc., Brattleboro, VT). The signal was recorded at 100 Hz with a photomultiplier (Nikon P600S) and a computerized acquisition system (40). The cell layer was perfused at 1 ml/min with the 500 nm free Ca$^{2+}$ intracellular buffer (see above) supplemented with 0.1 $\mu$g/ml rhodamine 123.

*Succinate Oxidation to CO$_2$ in Permeabilized INS-1 Cells—* INS-1 cells were seeded at 4 x $10^5$ cells/ml on 35-mm diameter dishes coated with A431 extracellular matrix as described above. Cells were maintained in glucose-free RPMI 1640 medium to subconfluence. Attached cells were then incubated in glucose- and glutamine-free RPMI 1640 medium plus 10 mM HEPES for 2 h at 37 °C, transferred to a thermostatted glass chamber, and permeabilized according to the procedure described above. Cells were then washed with the corresponding intracellular buffer adjusted to either 100 or 500 nm free Ca$^{2+}$ and preincubated for 10 min in that buffer. Succinate oxidation was initiated by replacing the buffer with 1 ml of the respective fresh ones containing 1 mM (23, 34) succinate (NEN Life Science Products; 0.1 $\mu$l/chamber). After a 1-h incubation at 37 °C in sealed chambers, 0.5 ml of 0.1 M HCl was added onto the cell layers to stop the reaction, and 1 ml of benzethonium hydroxide (Sigma) was injected into the bottom of the chamber to bind the CO$_2$ liberated by the reaction (41). Following an overnight incubation at room temperature, 14CO$_2$ production was measured in benzethonium extracted with 5 ml of EtOH and counted in an LS6500 liquid scintillation counter (Beckman Instruments).

**Statistical Analysis—** Where applicable, values are expressed as the mean ± S.E., and significance of difference was calculated by Student’s t-test for unpaired data. Traces without S.E. values are representative of at least three independent experiments.

**RESULTS**

*Insulin Secretion in Islets—* Rat pancreatic islets were maintained in culture for 2–4 days prior to the experiments. Stimulation of insulin secretion with 16.7 mM glucose for 10 min was repeated after a 10-min interval of perfusion at 2.8 mM glucose. This revealed that the secretory response was desensitized during the second stimulation, displaying ~50% reduction (Fig. 4A). The tricarboxylic acid cycle intermediate succinate, rendered cell-permeant by the ester binding of a methyl group (42), also produced a desensitization of the insulin exocytotic response with a pattern similar to that produced by glucose (Fig. 1B). Finally, KCl was used to raise [Ca$^{2+}$] by membrane depolarization (29, 17). Again, the second of two exposures to 20 mM KCl revealed a blunted insulin secretory response (Fig. 1C).

Mitochondrial Desensitization and Insulin Secretion
Primary Pancreatic Cells—Primary rat pancreatic cells were transiently transfected with the cDNA encoding mitochondrial targeted aequorin. Monitoring of \([Ca^{2+}]_m\) in these cells showed that 5 mM methyl succinate increased \([Ca^{2+}]_m\) during the first stimulation, but not during a second one repeated 5 min later (Fig. 2A). This desensitization was also observed by raising \([Ca^{2+}]_c\) through depolarization of the plasma membrane induced by 20 mM KCl (Fig. 2B). Contrary to clones stably expressing aequorin, the low expression levels after transient transfection (13) do not permit a reliable calibration since the total photon emission was 10–20-fold less in the later case. Therefore, \([Ca^{2+}]_m\) is expressed as photons emitted per second.

ATP Generation, \([Ca^{2+}]_m\), and Insulin Secretion in INS-1 Cells—The insulin-secreting cell line INS-1 was stably transfected with mitochondrially targeted aequorin (INS-1/EK3) or with luciferase (INS-r3-LUC7), allowing the continuous measurement of \([Ca^{2+}]_m\) or [ATP], respectively, in living perifused cells. The simultaneous monitoring of \([Ca^{2+}]_m\) and insulin secretion demonstrated that both parameters exhibited an attenuated response when 5 mM methyl succinate was added to the perifusion 5 min after the first stimulation (Fig. 3, B and C, respectively). The \([Ca^{2+}]_m\) desensitization was not due to aequorin consumption or deleterious effects on the cells, as the \([Ca^{2+}]_m\) response to methyl succinate was completely restored after an interval of 30 min between the two pulses (Fig. 3D).

The addition of 5 mM methyl succinate to INS-1 cells produced an increase in cytosolic ATP, and the same rise could be elicited 5 min later to the same extent during a second exposure to methyl succinate without any significant desensitization (Fig. 3A). Additional time points for the \([Ca^{2+}]_m\) increases and recovery of the secretory responses have already been documented using glucose as a stimulus (23). Moreover, glucose, which also increases cytosolic ATP levels (36), did not exhibit any desensitization using the protocol of Fig. 2A. The ATP response to 12.8 mM glucose was \(+23.3 \pm 2.0\) and \(+24.3 \pm 2.2\%\) during the first and second applications, respectively (not significant, \(n = 4\)).

\([Ca^{2+}]_m\) in Permeabilized INS-1 Cells—The aequorin-expressing cells were then permeabilized with Staphylococcus a-toxin, which forms very small holes (2–3 nm diameter) in the plasma membrane (38, 8). In this preparation, the cytosolic composition and hence the mitochondrial environment can be controlled. The permeabilized cells were perifused with an intracellular type buffer containing a permissive free \(Ca^{2+}\) concentration of 500 nM and 10 mM ATP. The first addition of 1

![Fig. 1. Desensitization of insulin secretion in rat islets following repeated stimulation.](image-url) Rat pancreatic islets were isolated and kept in culture for 2–4 days before insulin secretion experiments in a perifusion system. Stimulation with 16.7 mM Glc for 10 min was repeated after a 10-min interval (A). Using the same protocol, the effect of a 5 mM concentration of the mitochondrial substrate methyl succinate (met-Suc) is shown in B. Insulin exocytosis was also stimulated by depolarizing the cells with 20 mM KCl (C). Values are the mean ± S.E. (n = 4).

![Fig. 2. Desensitization of \([Ca^{2+}]_m\) increases in rat islet cells.](image-url) Rat islet cells were transiently transfected with the cDNA encoding mitochondrially targeted aequorin and used 3 days later. Cells were perifused at 37 °C with HEPES-balanced Krebs-Ringer bicarbonate buffer and exposed for 5 min to 5 mM methyl succinate (met-Suc) (A) or 20 mM KCl (B). Stimulation was repeated after a 5-min interval. The traces are representative of at least three independent experiments.
mM succinate induced a large transient peak in $[\text{Ca}^{2+}]_m$, but the second pulse 5 min later was ineffective (Fig. 4A). The desensitization phenomenon was also observed with $\alpha$-glycerophosphate, which transfers reducing equivalents from the cytosol to the same site (complex II) in the electron transport chain as succinate (Fig. 4B). Glycerophosphate has been shown to produce ATP in isolated islet mitochondria (43). More important, using 5-min intervals, succinate desensitized the effect of $\alpha$-glycerophosphate on $[\text{Ca}^{2+}]_m$ and vice versa (Fig. 4C and D). This latter effect shows that the desensitization mechanism appears to be located downstream of the oxidation of FADH$_2$. It should be noted that when Ca$^{2+}$ was substituted with the Ca$^{2+}$ surrogate Sr$^{2+}$ in the intracellular type buffer, a very similar desensitization of the mitochondrial [Sr$^{2+}$] increase was observed upon repeated succinate stimulation.\(^2\) As for intact cells, the desensitization was not an irreversible process due to a toxic effect or to the loss of functional aequorin since resensitization was observed after 30 min using either succinate or $\alpha$-glycerophosphate (Fig. 5, A and B, respectively).

To examine whether a $[\text{Ca}^{2+}]_m$ increase per se causes desensitization to subsequent challenges, the free Ca$^{2+}$ concentration of the buffer was varied. To this end, the extramitochondrial $[\text{Ca}^{2+}]_m$ in permeabilized cells was kept at permissive 500 nM levels or raised to 1.3 $\mu$M. The stimulation for 5 min with 1.3 $\mu$M Ca$^{2+}$ induced a first transient peak in $[\text{Ca}^{2+}]_m$ up to 1.7 $\mu$M,

\(^2\) P. Maechler, E. D. Kennedy, and C. B. Wollheim, unpublished observations.
followed by a second phase that tended to stabilize to the level of extramitochondrial Ca\(^{2+}\). The second exposure to 1.3 mM Ca\(^{2+}\) showed a complete desensitization of the first transient increase in [Ca\(^{2+}\)]\(_{m}\) above the equilibration value between the two compartments (Fig. 6). Imposing the same experimental protocol except for a shortening of the exposures to 1.3 mM Ca\(^{2+}\) to 30 s instead of 5 min did not result in any desensitization of the [Ca\(^{2+}\)]\(_{m}\). The latter observation is in agreement with results reported for permeabilized HeLa cells expressing mitochondrial aequorin (44).

**Effect of Inhibitors of the Electron Transport Chain on [Ca\(^{2+}\)]\(_{m}\) in Permeabilized INS-1 Cells**—Succinate dehydrogenase generates FADH\(_{2}\), with the subsequent transfer of electrons to complex II of the electron transport chain (45). In permeabilized cells, the effect of succinate on the increase in [Ca\(^{2+}\)]\(_{m}\) was not affected by the presence of 100 mM rotenone, which blocks complex I of the respiratory chain (Fig. 7A). On the other hand, the succinate-induced [Ca\(^{2+}\)]\(_{m}\) increase was completely abolished by 10 mM antimycin A, an inhibitor of complex III (Fig. 7B). This suggests that the desensitization occurs between complex II and the uniporter, the latter mediating Ca\(^{2+}\) uptake in the mitochondria.

**Effect of Free Ca\(^{2+}\) Concentration on DC\(_{m}\) and [Ca\(^{2+}\)]\(_{m}\) in Permeabilized INS-1 Cells**—Ca\(^{2+}\) influx into the mitochondria through the uniporter is driven by the hyperpolarization of the mitochondrial membrane under conditions of permissive [Ca\(^{2+}\)]\(_{m}\). The hyperpolarization occurs by the transfer of reducing equivalents to the electron transport chain and the resulting extrusion of protons. We next tried to discriminate between the respiratory chain and the uniporter as the site of desensitization. For this purpose, we studied the effect of succinate on [Ca\(^{2+}\)]\(_{m}\) in permeabilized cells perifused with nonpermissive (resting) or permissive free [Ca\(^{2+}\)] (100 and 500 nM, respectively). Under both conditions, succinate was efficient in hyperpolarizing the mitochondrial membrane (Fig. 8, A and B). The dissipation of the proton gradient by carbonyl cyanide p-triflu-
The effects of repeated 5-min exposures to 1.3 mM succinate on 
mitochondrial aequorin were permeabilized with α-toxin and perfused with a buffer containing 500 nM free Ca^{2+} and 10 mM ATP. The effects of repeated 5-min exposures to 1.3 mM succinate (final concentration) in the buffer on [Ca^{2+}]_m were tested with a 5-min interval. The trace is representative of four independent experiments.

**Effect of Extramitochondrial Ca^{2+} on Succinate Oxidation to CO_{2} in Permeabilized INS-1 Cells**—The hyperpolarizing action of succinate on ΔΨ_m is catalyzed by succinate dehydrogenase, a Ca^{2+}-independent enzyme (45). By contrast, CO_{2} formation from succinate requires a complete turn of the tricarboxylic acid cycle, which involves the two Ca^{2+}-sensitive enzymes NAD-isocitrate dehydrogenase and α-ketoglutarate dehydrogenase (15). As shown in Fig. 9, [2,3-14C]succinate oxidation to 14CO_{2} was stimulated 4-fold (p < 0.01) by an increase in the extramitochondrial [Ca^{2+}] from 100 to 500 nM in the permeabilized INS-1 cells.

**DISCUSSION**

Mitochondria play a key role in the metabolism-secretion coupling of the pancreatic beta cell (9–11, 13, 20). Evidence for a desensitization of this organelle is presented here, and the phenomenon may account for the documented desensitization of stimulated insulin secretion observed in pancreatic beta cells (1, 4, 5) and derived cell lines (23, 46). In the present study, repeated exposures of rat islets to stimulatory glucose concentrations led to attenuated insulin exocytosis. This desensitized secretory response was also observed with the mitochondrial substrate methyl succinate or with KCl-induced depolarization of the plasma membrane. In primary islet cells, desensitization of the mitochondria was indicated by the impaired rise of [Ca^{2+}]_m during the second exposure to methyl succinate or high potassium. This suggests that desensitization of Ca^{2+} entry into the mitochondria can be evoked by either tricarboxylic acid cycle intermediates or simply by increasing the [Ca^{2+}]_m. Nevertheless, to be considered a pure mitochondrial effect, the latter condition implies that the [Ca^{2+}]_m increase would reach the same value during the second exposure to KCl or at least a level well above the threshold of the uniporter (400 μM) (21, 16). Although desensitization of the [Ca^{2+}]_m response to KCl occurs in INS-1 cells, it still attains micromolar concentrations during the second pulse (23). The [Ca^{2+}]_m reduction is less marked than that of [Ca^{2+}]_m, and therefore probably plays only a minor role in the mitochondrial desensitization. In intact INS-1 cells stimulated with methyl succinate, the blunted insulin secretion correlated with an inhibited increase in [Ca^{2+}]_m upon a second exposure. In contrast, methyl succinate-induced generation of ATP, reflecting the activation of oxidative phosphorylation, did not display any desensitization, as demonstrated in luciferase-expressing INS-1 cells. The cellular responses to glucose are also desensitized with respect to [Ca^{2+}]_m and insulin secretion (23), but not in terms of ATP generation (see Results). This dichotomy between two mitochondrial parameters, [Ca^{2+}]_m and ATP generation, can be explained by...
reduced Ca$^{2+}$ uptake into the mitochondrial matrix, despite a fully activated respiratory chain. To investigate the underlying mechanism, we have used permeabilized cells to clamp extra-mitochondrial [Ca$^{2+}$] at a fixed permissive level of 500 nM. This was chosen to ascertain Ca$^{2+}$ uptake by the uniporter (16). Under these conditions, the succinate-induced increase in [Ca$^{2+}$]$_{m}$ was completely desensitized during the second stimulation. This inhibitory effect takes place downstream of complex II and is apparently not due to altered transport of succinate into the mitochondria. Indeed, the desensitizing effect of succinate could be reproduced with $\alpha$-glycerophosphate. This latter compound transfers reducing equivalents from the glycolytic intermediate dihydroxyacetone phosphate to the same complex II of the electron transport chain without being transported into the mitochondrial matrix (47). Thus, the desensitization evoked by both of the FADH$_2$-producing substances (succinate and $\alpha$-glycerophosphate) is very similar, and a common mode of action is underscored by a clear cross-desensitization. In addition, succinate-induced increases in [Ca$^{2+}$]$_{m}$ were blocked by inhibiting complex III with antimycin A, but not by rotenone, which blocks complex I. We therefore conclude that the site of desensitization is located downstream of complex II either in the electron transport chain or at the uniporter through which Ca$^{2+}$ flows into the mitochondria. The desensitization does not appear to be due to inhibition of the respiratory chain, the activation of which was not impaired. This is demonstrated by the hyperpolarization of $\Delta \Psi_m$ by succinate at 100 nM Ca$^{2+}$, a normal rise in [Ca$^{2+}$]$_{m}$ can be induced at permissive [Ca$^{2+}$] 5 min later with the same substance without any desensitization (C). In D, $\Delta \Psi_m$ was hyperpolarized during both exposures to 1 mM succinate in attached permeabilized INS-1 cells perfused with the 500 nM free Ca$^{2+}$ intracellular buffer. The traces are representative of at least three independent experiments. FCCP, carbonyl cyanide p-trifluoromethoxyphenylhydrazone; $\alpha$. $u.$, arbitrary units.
inhibitory effect of Ca\(^{2+}\) alone. KCl (20 mM) evokes increases in [Ca\(^{2+}\)]\(_{\text{m}}\) up to 2 \(\mu\)M (23). In permeabilized cells, direct applications of Ca\(^{2+}\) in this concentration range clearly caused desensitization of the [Ca\(^{2+}\)]\(_{\text{m}}\) response to the second pulse. This applies to the transient [Ca\(^{2+}\)]\(_{\text{m}}\) increase, but not to the equilibration of the ion between the extra- and intramitochondrial compartments, which suggests two independent pathways for mitochondrial Ca\(^{2+}\) uptake. Moreover, the desensitization requires a complete activation involving a new steady state. Indeed, very short applications (<1 min) of Ca\(^{2+}\) in the micromolar range do not lead to desensitization of [Ca\(^{2+}\)]\(_{\text{m}}\) responses during a second stimulation in permeabilized cells (44). Although the molecular nature of the mitochondrial Ca\(^{2+}\) uniporter has not been identified, it appears to have properties similar to those of Ca\(^{2+}\) channels of the plasma membrane (48). It may therefore be speculated that the desensitization evoked by an increase in [Ca\(^{2+}\)]\(_{\text{m}}\) could involve a mechanism similar to that described for L-type Ca\(^{2+}\) channels (49, 50). Such Ca\(^{2+}\) channel desensitization has also been reported in insulin-secreting cells (51). It is conceivable that the high frequency of the [Ca\(^{2+}\)] oscillations (two to five/min) observed in glucose-stimulated beta cells (12, 17, 52) serves to prevent desensitization of mitochondrial metabolism. It may be important to optimize the activity of the Ca\(^{2+}\)-sensitive dehydrogenases of the mitochondria to ensure the continuous production of metabolic coupling factors. We show here that succinate oxidation, reflecting tricarboxylic acid cycle activity, is stimulated by extramitochondrial [Ca\(^{2+}\)] in the physiological concentration range (500 nM). Such an effect was previously reported for the oxidation of pyruvate and its conversion to citrate (20).

The consensus model of metabolism-secretion coupling in the beta cell attributes a key role to ATP produced by the mitochondria (9, 10, 31). However, as clearly demonstrated by repeated stimulation with methyl succinate, ATP generation is not sufficient for the triggering of insulin secretion. Hence, in intact INS-1 cells, ATP production was preserved in the face of blunted [Ca\(^{2+}\)]\(_{\text{m}}\) and secretory responses during the second application of methyl succinate. This will result in diminished activation of the mitochondrial Ca\(^{2+}\)-sensitive dehydrogenases (15), the stimulation of which is required for full activation of the mitochondrial metabolism. An unidentified mitochondrial factor, distinct from ATP, has been proposed to participate in the triggering of insulin exocytosis (13). Its generation requires both a rise in [Ca\(^{2+}\)]\(_{\text{m}}\) and the provision of carbons to the tricarboxylic acid cycle (13). Thus, we speculate that during the desensitization of the beta cell, despite normal ATP generation, this mitochondrial factor is missing due to insufficient elevation of [Ca\(^{2+}\)]\(_{\text{m}}\). As a consequence of deficient generation of coupling factors, insulin secretion is impaired. The nature of the coupling factors of mitochondrial origin remains to be established.

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REFERENCES

**FIG. 9.** Effect of extramitochondrial Ca\(^{2+}\) (100 or 500 nM) on [2,3-\(^{14}\)C]succinate oxidation to \(^{14}\)CO\(_2\) in a-toxin-permeabilized INS-1 cells. Cells were first permeabilized for 10 min and then equilibrated for another 10-min period in the corresponding intracellular buffers prior to a 1-h stimulation with 1 mM [2,3-\(^{14}\)C]succinate (0.1 \(\mu\)Ci/chamber). Values are the mean \(\pm\) S.E. (\(n = 3\)) of one experiment representative of four independent experiments.
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