Measurements of membrane potential, transmembrane $^{45}$Ca fluxes, cytoplasmic free Ca$^{2+}$ concentration and insulin release by transplantable rat insulinoma cells maintained in tissue culture

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Summary Regulation of insulin release, membrane potential, transmembrane $^{45}$Ca fluxes and cytoplasmic free Ca$^{2+}$ concentration, [Ca$^{2+}$], was examined using suspensions of transplantable NEDH rat insulinoma cells previously cultivated for 2-3 days to eliminate necrotic tumour cells and counter prior hypoglycaemia. Insulinoma cells displayed a resting [Ca$^{2+}$] of 94±8 nM (n=17) and released 104±15 ng insulin 10$^{-6}$ cells (n=7) during 60 min incubations with uptake of 2.7±0.2 mmol $^{45}$Ca 10$^{-6}$ cells (n=7). High concentrations of glucose did not affect membrane potential, transmembrane $^{45}$Ca fluxes, [Ca$^{2+}$], or insulin release by insulinoma cells. K$^{+}$ at 25 mM depolarised the plasma membrane, induced a small increase in $^{45}$Ca influx and increased [Ca$^{2+}$], by 65%. This modest action was not associated with demonstrable effects on Ca$^{2+}$ uptake and insulin release. The effect of 25 mM K$^{+}$ on [Ca$^{2+}$], was counteracted by D-600, but this blocker of voltage-activated Ca$^{2+}$ channels and verapamil lacked effects on transmembrane $^{45}$Ca fluxes and insulin release. The Ca$^{2+}$-calmodulin antagonist, trifluoroperazine, was also without effect on $^{45}$Ca fluxes and insulin release. Ca$^{2+}$ ionophore ionomycin increased [Ca$^{2+}$], whereas A23187 and X537A did not affect transmembrane $^{45}$Ca fluxes. Moreover, insulin release was independent of extracellular Ca$^{2+}$ over the range 0-20 mM despite marked affects on transmembrane $^{45}$Ca fluxes and a greater than 4-fold change of [Ca$^{2+}$]. Dibutyryl cyclic AMP increased insulin release by 55%, without affecting transmembrane $^{45}$Ca fluxes or [Ca$^{2+}$]. The phosphodiesterase inhibitor, theophylline, also enhanced insulin release by 10-36%, with no change of Ca$^{2+}$ uptake. The effectiveness of theophylline was independent of extracellular Ca$^{2+}$ over the range 0-10 mM. These results indicate that inappropriate Ca$^{2+}$ regulation is a key pathogenic feature underlying the inappropriate insulin secretion of rat insulinoma cells.

Recent years have witnessed a considerable interest in the development and exploitation of transplantable insulinomas in small laboratory animals for studies of the physiology and pathophysiology of insulin secretion (Grillo et al, 1967; Chick et al., 1977, 1980; Hirayama et al., 1979; Hanahan, 1985). The most commonly employed tumour is the serially transplantable radiation-induced NEDH rat insulinoma (Chick et al., 1977) which exhibits rapid growth rate, giving rise to large vascularised tumours comprised of well granulated insulin-containing cells with only trace amounts of other regulatory peptides (O’Hare et al., 1985; Conlon et al., 1986). This tumour also has parented the clonal RINm5F cell line (Gazdar et al., 1980), although the insulin-secretory properties of this selected daughter clone more closely resemble pancreatic β-cells than the original tumour (Flatt et al., 1987a).

Syngeneic transplantation of small fragments of the NEDH rat insulinoma consistently results in hyperphagia, loss of diurnal rhythms of insulin-glucose homeostasis, progressive hyperinsulinemia and hypoglycaemia which without surgical or drug intervention results in neuroglycopenic coma (Flatt et al., 1986, 1987b, c, d). Since uncontrolled insulin secretion in the face of hypoglycaemia is the cardinal feature of insulinoma (Marks & Rose, 1981), recent studies have focused on the regulation of tumour insulin secretion in relation to the underlying secretory defect (Swanson-Flatt & Flatt, 1987, 1988a,b). These studies have shown that whereas rat insulinoma cells respond to agents which affect insulin secretion in pancreatic β-cells through the adenylate cyclase-cyclic AMP system, responsiveness to glucose and substances which normally modulate secretion by alterations of transmembrane Ca$^{2+}$ fluxes is severely compromized. To further evaluate this issue, the present study has examined the role of extracellular Ca$^{2+}$ in insulin release from cultured rat insulinoma cells, and assessed changes in membrane potential, transmembrane $^{45}$Ca flux, cytoplasmic free Ca$^{2+}$ concentration, [Ca$^{2+}$], and insulin release following exposure to nutrients and drugs with established effects on pancreatic β-cells.

Materials and methods

Animals and transplantation

Male inbred albino New England Deaconess Hospital (NEDH) rats from the colony at the University of Surrey carrying a serially transplantable radiation-induced insulinoma (Chick et al., 1977) were used at 14–17 weeks of age. The origin, transplantation and maintenance of these rats has been described elsewhere (Flatt et al., 1986).

Isolation and culture of tumour cells

Tumours were excised from the subcapsular site of hypoglycaemic insulinoma-bearing rats, and used to prepare tumour cell suspensions as previously described (Flatt et al., 1987a; Swanston-Flatt & Flatt, 1987). Isolated cells were cultured for 2–3 days at 37°C in a humidified atmosphere of 5% CO$_2$ in air. The culture medium was RPMI-1640 (Gibco Europe Ltd., Paisley, UK) containing 10% foetal calf serum, antibiotics (100 U ml$^{-1}$ penicillin and 0.1 mg ml$^{-1}$ streptomycin; Gibco Europe Ltd., Paisley, UK) and 11.1 mM glucose.

Measurements of insulin release and $^{45}$Ca uptake

Insulin release and $^{45}$Ca uptake studies were performed by incubating approximately 0.5 x 10$^6$ viable tumour cells in 100 μl modified Krebs Ringer buffer (pH 7.4) containing 20 mM N-2-hydroxyethylpiperazine-N’-2-ethanesulfonic acid (HEPES), 115 mM NaCl, 24 mM NaHCO$_3$, 4.7 mM KCl, 2.6 mM $^{45}$CaCl$_2$ (7.8 Ci mol$^{-1}$; Amerham International, Amersham, UK) 1.2 mM KH$_2$PO$_4$, 1.2 mM MgSO$_4$ and 5 mg ml$^{-1}$ bovine serum albumin. The buffer was supplemented with glucose and other test agents as indicated in the Tables and Figures. In experiments involving substantial
changes of extracellular Ca\(^{2+}\) concentration, control incubations were performed using choline chloride to correct for osmotic pressure changes.

The cells were incubated for 60 min at 37°C in polyethylene microfuge tubes (400 μl capacity) in triplicate. At the end of incubation, the cells were separated by centrifugation (Beckman microfuge type B, 15,000 g for 2 min) through an underlying layer of oil (200 μl of a mixture of dibutyryl-phosphodiester and dinonylphthalate, 10:3 vol/vol) into urea (20 μl, 6 M). Aliquots of supernatant buffer were stored at −20°C for insulin assay (Flatt & Bailey, 1981). The \(^{45}\)Ca content of the cell pellet in the cut tip of the tube was determined by liquid scintillation spectrometry following the addition of 2 ml Piccolfluor-15 (Packard Instruments Ltd., Caversham, UK). Samples of the labelled incubation medium (5 μl) were used as external standards in the counting procedure.

**Measurements of \(^{45}\)Ca efflux**
\(^{45}\)Ca efflux studies were performed using cultured tumour cells preloaded with \(^{45}\)Ca during incubation for 90 min in 100 μl of the Kreb's Ringer buffer supplemented with 2.6 mM \(^{45}\)Ca (360 Ci mmol\(^{-1}\)) and 20 mM glucose. After briefly washing in nonradioactive buffer, approximately 2 × 10\(^6\) tumour cells were transferred to 10 μl chambers and perfused at a constant rate of approximately 40 μl min\(^{-1}\) (Flatt et al., 1980a). The perfusate was collected over successive periods of 1 or 5 min, with inclusion of glucose and other test agents in the buffer from 35−75 min as indicated in Figures 1 and 2. Samples of the perfusate (15 μl) were mixed with 2 ml Piccolfluor-15 (Packard Instruments Ltd., Caversham, UK) and analysed for radioactivity by liquid scintillation counting. In each individual experiment, the \(^{45}\)Ca efflux rate (C.P.M. per minute) was expressed as a percentage of the mean value observed in the same experiment between the 31st and 36th minutes of perfusion.

**Measurements of cytoplasmic free Ca\(^{2+}\)  ion concentration**
Studies of cytoplasmic free Ca\(^{2+}\) concentration were performed using cultured tumour cells previously incubated for 45 min in Kreb's Ringer buffer containing 5 μM quin-2-AM (Sigma Chemical Co. Ltd., Poole, Dorset). The loading was ~3.2 nmol quin-210\(^{-10}\) cells, as judged from calculations based on fluorescence maximum and extracellular quin-2 values at time zero, assuming that 1 mg dry weight corresponds to 3.6 × 10\(^6\) cells (Lenmark, 1974). After briefly washing, the cells were resuspended in 1.5 ml buffer and transferred to the cuvette. Fluorescence was measured with excitation and emission wavelengths set at 339 and 492 nm, respectively. Calibration was performed as previously described (Beaven et al., 1984).

**Measurements of membrane potential**
Qualitative changes in membrane potential of cultured tumour cells were measured with the fluorescent dye, bis-oxonol (Rink et al., 1980; Molecular Probes, Junction City, Oregon, USA). Bis-oxonol, at a final concentration of 150 nM, was allowed to equilibrate with 1.5 ml Kreb's Ringer buffer before transferring the cells to the cuvette. Measurements were performed at excitation and emission wavelengths of 540 and 580 nm, respectively. In common with studies of [Ca\(^{2+}\)], fluorescence was determined at 37°C continuously during exposure to glucose and other test agents using an Aminco-Bowman spectrofluorometer, slightly modified to allow constant stirring in 1 cm polystyrene cuvettes.

**Statistical analysis**
Values are presented where appropriate as mean ± s.e.m. Statistical evaluation was performed using analysis of variance (Anova) and Student's paired or unpaired t-tests. Differences were considered to be significant for P < 0.05.

**Results**

**Insulin release and \(^{45}\)Ca uptake**
As shown in Table I, glucose did not modify insulin release or \(^{45}\)Ca uptake by rat insulinoma cells. Addition of theophylline or dibutyryl cyclic AMP enhanced insulin release by 19-52%, without affecting \(^{45}\)Ca uptake. Figure 1 shows \(^{45}\)Ca uptake and insulin release at different extracellular Ca\(^{2+}\) concentrations in the absence or presence of theophylline. The extent of \(^{45}\)Ca uptake increased with Ca\(^{2+}\) concentration, and was not saturable between 0-20.4 mm. Changes of extracellular Ca\(^{2+}\) over this range did not affect insulin release irrespective of the presence or absence of theophylline. However, theophylline increased insulin release by 10-30%, compared with control incubations performed at 0-10 mm Ca\(^{2+}\). Theophylline lacked effects on insulin release at 20.4 mm Ca\(^{2+}\), and was without effect on \(^{45}\)Ca uptake irrespective of Ca\(^{2+}\) concentration.

As shown in Table II, \(^{45}\)Ca uptake and insulin release by rat insulinoma cells were not affected by a depolarising concentration of K\(^+\). Furthermore, blockade of voltage-dependent Ca\(^{2+}\) channels using verapamil or D-600, addition of the Ca\(^{2+}\)-calmodulin antagonist, trifluoroperazine, or the calcium ionophore X537A failed to modify \(^{45}\)Ca uptake or insulin release. The calcium ionophore A23187 increased insulin output by 47%, without change of \(^{45}\)Ca uptake. DMSO which was used to disolve ionophores did not affect \(^{45}\)Ca uptake or insulin release at the final concentration of 0.1%.

**\(^{45}\)Ca efflux**
Whereas the measurements of \(^{45}\)Ca uptake give an indication of the balance between increased influx and the efflux of radioactive \(^{45}\)Ca by the cell, evaluation of \(^{45}\)Ca efflux provides a highly sensitive and continuous measurement of unidirectional Ca\(^{2+}\) flux. As shown in Figure 2 (A-C) glucose, dibutyryl cyclic AMP, A23187 or verapamil lacked significant effects on \(^{45}\)Ca efflux from preloaded rat insulinoma cells. In contrast to its ineffectiveness in \(^{45}\)Ca uptake studies, 25 mm K\(^+\) caused a slight transient but significant increase of \(^{45}\)Ca efflux (Figure 2A). Changes in the medium concentration of Ca\(^{2+}\) also significantly affected \(^{45}\)Ca efflux (Figure 3 A-B). Addition or removal of Ca\(^{2+}\) from the

**Table I** Effects of glucose, theophylline and dibutyryl cyclic AMP on \(^{45}\)Ca uptake and insulin release by rat insulinoma cells

| Glucose (mM) | Additions | \(^{45}\)Ca uptake (nmol 10\(^{-6}\) cells h\(^{-1}\)) | Insulin release (ng 10\(^{-6}\) cells h\(^{-1}\)) |
|-------------|-----------|----------------|------------------|
| 0           | —         | 2.65 ± 0.22    | 104 ± 15         |
| 1.4         | —         | 2.84 ± 0.58    | 119 ± 30         |
| 16.7        | —         | 3.67 ± 0.94    | 102 ± 19         |
| 16.7        | theophylline (5) | 3.44 ± 0.68 | 134 ± 22         |
| 16.7        | db cyclic AMP (2.5) | 3.35 ± 0.46 | 158 ± 11         |

*Values are mean ± s.e.m. of 7 experiments; *P < 0.02 (at least) compared with 0 mM glucose or 16.7 mM glucose.
perfusion medium transiently increased the 45Ca efflux rate from insulinoma cells.

**Cytoplasmic free Ca2+ concentration**

Measurements of cytoplasmic free Ca2+ ion concentration, [Ca2+]i, in rat insulinoma cells gave a resting [Ca2+]i of 94±8 nm (mean ±s.e.m., n = 17). As shown in Table III and the representative traces in Figure 4A-D, glucose did not affect [Ca2+]i, whereas exposure to 25 mM K+ resulted in a modest increase of [Ca2+]i by 65%. The effect of 25 mM K+ was counteracted by the voltage-dependent Ca2+ channel blocker D-600 (40% decrease of [Ca2+]i; Figure 4B). In other experiments (Figure 4C), dibutyryl cyclic AMP did not affect [Ca2+]i. This nucleotide itself induced a small degree of autofluorescence (Figure 4C, lower trace). The Ca2+ ionophore, ionomycin, increased [Ca2+]i, by 81% (Figure 4D).

The effects of manipulation of extracellular Ca2+ concentration on [Ca2+]i, are shown in Figures 5 and 6. Increasing extracellular Ca2+ stepwise from 0 mM to 0.26 mM, and through to 20.4 mM resulted in successive increases in [Ca2+]i (each step at least P<0.05, except between 5.1 mM and 10.2 mM which did not achieve significance). [Ca2+]i appeared not to be saturable over the range 0–20.4 mM extracellular Ca2+.

**Membrane potential**

Relative changes in membrane potential in rat insulinoma cells were measured using the fluorescent dye bis-oxonol. As shown in Figure 7, glucose was without effect on the membrane potential of insulinoma cells. Exposure to 25 mM K+ resulted in depolarisation of the plasma membrane.

**Discussion**

Considerable evidence supports a key role of Ca2+ as a regulator of insulin secretion from the pancreatic β-cells (Malaisse et al., 1978a; Wollheim & Sharp, 1981; Henquin & Meissner, 1984; Hellman & Gylfe, 1986). Thus glucose and many other secretagogues are believed to regulate insulin release by controlling the concentration of free Ca2+ ions in a stimulatory cytoplasmic pool. This is achieved through effects on Ca2+ fluxes at the plasma membrane and intracellular Ca2+ sequestration by organelles such as mitochondria and endoplasmic reticulum. Elevation of intracellular cyclic AMP also triggers insulin release in the presence of extracellular Ca2+, and it is generally held that one important action of cyclic AMP in addition to activation of protein kinases concerns sensitization of the exocytotic mechanism to cytoplasmic Ca2+ (Wollheim & Sharp, 1981; Hellman & Gylfe, 1986). In the present study, we have demonstrated that cultured rat insulinoma cells exhibit profound irregularities in the regulation of transmembrane Ca2+ fluxes, insulin release and [Ca2+]i, measured using the fluorescent indicator quin-2.

It is now well established that in pancreatic β-cells glucose triggers a network of interrelated metabolic and ionic events which lead to depolarisation of the plasma membrane, Ca2+ influx, elevation of [Ca2+]i, and insulin release (Malaisse et al., 1978a; Wollheim & Sharp, 1981; Henquin & Meissner, 1984; Hellman & Gylfe, 1986). In accordance with previous studies with the Surrey insulinoma subline (Flatt et al., 1987b; Swanston-Flatt & Flatt, 1987), short-term cultured rat insulinoma cells did not respond to glucose with

![Graph](image)

**Figure 1** Effects of extracellular Ca2+ concentration on insulin release and 45Ca uptake by rat insulinoma cells. Incubations were performed at 11.1 mM glucose in the absence (○) or presence (□) of 5 mM theophylline. Values are mean ±s.e.m. of 7 experiments. *P<0.05 (at least) compared with insulin release in the absence of theophylline. 45Ca uptake was consistently increased at least P<0.05 by stepwise increments of extracellular Ca2+ over the range from 0 mM (plus 1 mM EGTA) to 0.26 mM, 5.1 mM, 10.2 mM and 20.4 mM. Modulation of extracellular Ca2+ was without effect on insulin release. It was checked in control experiments using choline chloride that osmotic pressure changes did not affect insulin release.

| Additions (mm) | 45Ca uptake (nmol 10⁶ cells h⁻¹) | Insulin release (ng 10⁶ cells h⁻¹) |
|---------------|---------------------------------|----------------------------------|
| None (control)| 2.43 ± 0.59                     | 139 ± 12                         |
| K⁺ (25)       | 3.07 ± 0.54                     | 145 ± 10                         |
| Verapamil (0.05) | 3.64 ± 0.93                  | 142 ± 13                         |
| D-600 (0.05)  | 3.42 ± 0.79                     | 128 ± 9                          |
| Trifluoperazine (0.025) | 3.47 ± 1.08                | 134 ± 7                          |
| X537A (0.04)  | 3.97 ± 1.10                     | 144 ± 12                         |
| A23187 (0.02) | 5.90 ± 2.40                     | 204 ± 23                         |
| DMSO (0.1% control) | 3.59 ± 1.08                | 147 ± 13                         |

All incubations were performed in the presence of 11.1 mM glucose. Values are mean ±s.e.m. of 8 experiments. *P<0.01 compared with controls.
Figure 2 Effects of glucose, K+, dibutyryl cyclic AMP, A23187 and verapamil on $^{45}$Ca efflux from rat insulinoma cells. Experiments were performed in the parallel channels of a perfusion apparatus with rat insulinoma cells preloaded with $^{45}$Ca. The cells were perfused with buffer containing 1.4mM glucose, with exposure to the following test agents for 35-70min: Panel a, 16.7mM glucose (●) or 25mM K+ (▲); Panel b, 2.5mM dibutyryl cyclic AMP (●); Panel c, 20μM A23187 (●) or 50μM verapamil (▲). Open symbols (○) in each of the panels refer to control perifusions. Values are mean ± s.e.m. of 4-9 experiments. 25mM K+ significantly increased $^{45}$Ca efflux by 45min (ANOVA; $P<0.001$).

Figure 3 Effects of extracellular Ca$^{2+}$ concentration on $^{45}$Ca efflux from rat insulinoma cells. Experiments were performed in the parallel channels of a perfusion apparatus with rat insulinoma cells preloaded $^{45}$Ca. The cells were perfused with buffer containing 1.4mM glucose, with exposure to ionic manipulations for 35-70min: Panel a, change of Ca$^{2+}$ from 2.56mM to 0mM plus 1mM EGTA; Panel b, change of Ca$^{2+}$ from 0mM plus 1mM EGTA to 10.2mM Ca$^{2+}$. Open symbols (○) in each of the panels refer to control perifusions. Values are mean ± s.e.m. of 3-4 experiments. Ca$^{2+}$ removal or addition significantly increased $^{45}$Ca efflux by 45min (ANOVA; $P<0.01$ and $P<0.02$, respectively).
Table III  Effects of glucose, K+, D-600 and dibutyryl cyclic AMP on [Ca²⁺]i in rat insulinoma cells.

| Control | Test         | Cytoplasmic Ca²⁺ concentration (nM) |
|---------|--------------|-----------------------------------|
|         |              | Control (a)                        | Test (b)                          | Test – Control (b) |
| 1 Rest  | 20 mM glucose| 134 ± 11 (5)                       | 127 ± 11 (5)                      | -7 ± 10 (5)        |
| 2 Rest  | 25 mM K⁺     | 110 ± 11 (9)                       | 182 ± 14 (9)                      | +72 ± 16 (9)*      |
| 3 25 mM K⁺ | 50 μM D-600  | 179 ± 12 (8)                       | 108 ± 12 (8)                      | -71 ± 10 (8)*      |
| 4 Rest  | 2.5 mM db-cAMP| 76 ± 18 (4)                       | 78 ± 18 (4)                       | +2 ± 4 (4)         |
| 5 Rest  | 2 μM ionomycin| 141 ± 76 (2)                      | 292 ± 174 (2)                      | +151 ± 97 (2)      |

Values are mean ± s.e.m. of the number of experiments indicated in parenthesis. *P<0.01; *P<0.001 compared with appropriate control. The control buffer contained 0 mM glucose (experiments 1 and 5), 11.1 mM glucose (experiment 4) or 20 mM glucose (experiments 2 and 3). Cytoplasmic Ca²⁺ concentrations, calculated from stable fluorescence values before and after the addition of test agent, were determined from experiments like those shown in Figure 4.

Figure 4  Effects of glucose, K⁺, D-600, dibutyryl cyclic AMP and ionomycin on [Ca²⁺]i, in rat insulinoma cells. Fluorescent traces were obtained from rat insulinoma cells loaded with quin-2. Traces a–c are typical of experiments repeated 4–9 times. Trace d is representative of 2 experiments. The lower traces in a and c indicate autofluorescence due to the test substances per se. Approximate cytoplasmic Ca²⁺ concentrations are indicated as well as the additions of test agents.

Figure 5  Effect of extracellular Ca²⁺ concentration on [Ca²⁺]i in rat insulinoma cells. Fluorescent traces were obtained from rat insulinoma cells loaded with quin-2. The trace shown is typical of experiments repeated 4 times. Approximate cytoplasmic Ca²⁺ concentrations are given as well as the increments in extracellular Ca²⁺.
Figure 6 Relationship between extracellular Ca\(^{2+}\) concentration and [Ca\(^{2+}\)]\(_i\), in rat insulinoma cells. Cytoplasmic Ca\(^{2+}\) concentrations, calculated from stable fluorescence values after each addition of Ca\(^{2+}\), were derived from experiments like that shown in Figure 5. Values are mean ± s.e.m. of 4 experiments. Each stepwise increment in extracellular Ca\(^{2+}\), except between 5.1-10.2 mM, resulted in a significant increase in [Ca\(^{2+}\)]\(_i\), (P<0.05).

1988b) draw attention to possible irregularities in the Ca\(^{2+}\) channels of insulinoma cells.

The present study did not reveal a close correlation between increased concentrations of extracellular Ca\(^{2+}\) and insulin release by rat insulinoma cells. Thus whereas basal and nutrient-induced insulin secretion from rat pancreatic \(\beta\)-cells displays a marked Ca\(^{2+}\) dependence (Devi et al., 1977; Malaisse et al., 1978b), insulin release by insulinoma cells was not affected by depletion of Ca\(^{2+}\) from the medium with addition of the Ca\(^{2+}\)-chelator EGTA, or by increasing extracellular Ca\(^{2+}\) to 20.4 mM with or without osmotic compensation. Manipulations of extracellular Ca\(^{2+}\) were accompanied by dramatic changes in transmembrane \(^{45}\)Ca fluxes and [Ca\(^{2+}\)]\(_i\). The latter increased from 53±5 nm to 231±37 nm (n=4) in response to stepwise increments in extracellular Ca\(^{2+}\) over the range 0–20.4 mM. These observations indicate a disturbed sensitivity of the secretory process to Ca\(^{2+}\) in rat insulinoma cells, and accord with the general view that lack of a physiological Ca\(^{2+}\) response is a common determinant for the inappropiate functions of neoplastic cell types (Swierenga et al., 1980; Durham & Walton, 1982; Ralph, 1983). Such behaviour has been interpreted to indicate a paramount change in some key Ca\(^{2+}\)-regulated control mechanism (Swierenga et al., 1980; Ralph, 1983), and in insulinoma cells it is associated with the lack of effect of the Ca\(^{2+}\)-calmodulin antagonist, trifluoroperazine, on insulin release. However, the possibility cannot be ruled out that insulinoma cells exhibit exquisite sensitivity to Ca\(^{2+}\) such that insulin release is already maximally stimulated at extraordinary low [Ca\(^{2+}\)].

The ineffectiveness of the Ca\(^{2+}\) ionophore A23187 on transmembrane \(^{45}\)Ca fluxes in rat insulinoma cells parallels similar observations obtained with other neoplastic cell types (Cittadini et al., 1981). Since the present measurements of [Ca\(^{2+}\)]\(_i\) are slightly lower than reported for pancreatic \(\beta\)-cells (Rorsman et al., 1984; Rorsman & Abrahamsson, 1985; Abrahamsson et al., 1985), the phenomenon cannot be attributed to a high permeability of the cancer cell plasma membrane to external Ca\(^{2+}\). Ionophore X537A was also without effect on \(^{45}\)Ca uptake of rat insulinoma cells, although the Ca\(^{2+}\) ionophore ionomycin produced a prompt rise of [Ca\(^{2+}\)]\(_i\) in these cells. These combined observations may be taken to indicate that tumour cells in general have already acquired features similar to those induced by certain ionophores (see Cittadini et al., 1981). The increase of insulin release by A23187 in these circumstances may reflect an ability of calcium ionophores to promote cyclic AMP production in insulin-secreting cells (Hellman, 1975).

Previous in vivo and in vitro studies of rat insulinoma cells have drawn attention to the ability of agents which affect the adenylate cyclase-cyclic AMP system to modulate tumour insulin secretion (Flatt et al., 1987b; Swanston-Flatt & Flatt, 1988a). Consistent with this view, dibutyryl cyclic AMP and the phosphodiesterase inhibitor theophylline increased insulin release from rat insulinoma cells in the present study. The action of dibutyryl cyclic AMP did not involve modification of transmembrane \(^{45}\)Ca fluxes, and consistent with observations in pancreatic \(\beta\)-cells (Rorsman & Abrahamsson, 1985), [Ca\(^{2+}\)]\(_i\), was unchanged. Although the action of theophylline on pancreatic \(\beta\)-cells may involve mobilisation of cellular Ca\(^{2+}\) (Wollheim & Sharp, 1981; Hellman & Gylin, 1986), the changed Ca\(^{2+}\)-dependence of insulinoma cells clearly points to a primary effect of theophylline on intracellular cyclic AMP accumulation. Consistent with this view, theophylline-stimulation of insulin release was independent of extracellular Ca\(^{2+}\) over the range 0–10.2 mM, and the effect was not diminished by depletion of Ca\(^{2+}\) from the medium with addition of EGTA. This indicates that normal sensitivity to Ca\(^{2+}\) is not a prerequisite for cyclic AMP stimulated insulin release which may follow from activation of the operative microtubular-microfilamentous system in rat insulinoma cells (Swanston-Flatt & Flatt, 1988a). The fact that theophylline-induced insulin release was no longer demonstr-
able at 20.4 mM extracellular Ca2+ does not indicate an optimal [Ca2+]i for insulin release, as has been discussed in relation to Ca2+ -induced inhibition of nutrient-stimulated insulin release from pancreatic β-cells (Devis et al., 1977). High levels of Ca2+ may exert stabilising effects on the exocytotic mechanism triggered by cyclic AMP in insulinoma cells, or increased Ca2+ binding to intracellular sites in the plasma membrane may block insulin discharge as suggested by the inhibitory action of La3+ on rat insulinoma cells and pancreatic β-cells (Flatt et al., 1980a,b; Swanston-Flatt & Flatt, 1988a).

In conclusion, the present study has demonstrated that transplantable NEDH rat insulinoma cells exhibit marked abnormalities of insulin secretion associated with defective regulation of transmembrane Ca2+ fluxes and disturbances in both the control of and sensitivity to [Ca2+]i. Recently, it has been reported that inappropriate insulin release from three benign medullary-type human insulinomas was associated with disturbances in the regulation of transmembrane Ca2+ fluxes (Flatt et al., 1987e). Although some islet cell tumours may exhibit varying degrees of responsiveness to glucose or calcium ionophores (Henquin & Rose, 1981; Comi et al., 1986), the present observations indicate that inappropriate Ca2+ regulation is a common pathogenic feature underlying the inappropriate functions of certain types of insulinoma, and possibly other neoplastic cell types.

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