

**Therapeutic concentrations of antidepressants inhibit
pancreatic beta-cell function via mitochondrial complex
inhibition**

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5 **Therapeutic concentrations of antidepressants inhibit pancreatic beta-cell function via**
6 **mitochondrial complex inhibition**
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42 **Running title:** Antidepressant toxicity on beta-cells
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Abstract

Diabetes mellitus risk is increased by prolonged usage of antidepressants (ADs). Although various mechanisms are suggested for their diabetogenic potential, whether a direct effect of ADs on pancreatic β -cells is involved is unclear. We examined this idea for three ADs: paroxetine, clomipramine and, with particular emphasis, fluoxetine, on insulin secretion, mitochondrial function, cellular bioenergetics, K_{ATP} channel activity and caspase activity in murine and human cell-line models of pancreatic β -cells. Metabolic assays showed that these ADs decreased the redox, oxidative respiration and energetic potential of β -cells in a time and concentration dependent manner at concentrations (100 nM) well within their plasma therapeutic window. These effects were related to inhibition of mitochondrial complex I and III activity. Consistent with impaired mitochondrial function, lactate output was increased and insulin secretion decreased. Neither fluoxetine, antimycin nor rotenone could reactivate K_{ATP} channel activity blocked by glucose unlike the mitochondrial uncoupler, FCCP. Chronic, but not acute, AD increased oxidative stress and activated caspases, 3, 8 and 9. To conclude, paroxetine, clomipramine and fluoxetine were all found to be cytotoxic at therapeutic concentrations on pancreatic beta-cells; an action suggested to arise by inhibition of mitochondrial bioenergetics, oxidative stress and induction of apoptosis. These actions may help explain the diabetogenic potential of these ADs in humans.

Keywords: Antidepressant, Diabetes mellitus, beta-cell, mitochondria, cytotoxicity, fluoxetine.

Introduction

Antidepressants (ADs) play a key role in treating depression as well as other psychiatric disorders including anxiety and obsessive-compulsive disorders. However, clinical studies indicate that the tricyclic antidepressant (TCA) and serotonin selective reuptake inhibitor (SSRI) classes of AD are particularly associated with increased risks of hyperglycaemia and type 2 diabetes mellitus (T2D) (Lustman et al., 1997; Rubin et al., 2008; Brown et al., 2008; Derijks et al., 2008; Andersohn et al., 2009; Kivimäki et al., 2010; Khoza et al., 2011). Although weight gain is a common side effect of AD usage (Uguz et al., 2015) and a risk factor for T2D (Andersohn et al., 2009; Kivimäki et al., 2010;), when weight is controlled for, ADs still increase T2D risk (Rubin et al., 2008; Pan et al., 2012).

The effects of ADs on glucose homeostasis have been widely studied in rodents. The SSRI fluoxetine when administered acutely produces hyperglycaemia without hyperinsulinaemia (Yamada et al., 1999; Gomez et al., 2001), actions suggestive of a direct inhibition of insulin secretion. Similar trends are observed with clomipramine, a TCA, which also produces acute hyperglycaemia without hyperinsulinaemia in rodents (Sugimoto et al., 2003; Gomez et al., 2001) as well as in humans (Lustman et al., 1997; McIntyre et al., 2006). In murine models, intraperitoneal administration of fluoxetine or clomipramine produces maximal hyperglycaemia within the first 30-90 minutes of injection (Yamada et al., 1999; Sugimoto et al., 2003; Gomez et al., 2001); observations that also indicate a direct action of these drugs on the β -cell, an idea further supported by their ability to inhibit insulin secretion from isolated pancreatic β -cells (Antoine et al., 2004; Cataldo et al., 2015).

The mechanism of glucose stimulus-secretion coupling of insulin from the pancreatic β -cell is well understood, with a key role played by oxidative metabolism (Ashcroft et al., 1994; Maechler 2002). Clearly, any compound that interferes with mitochondrial bioenergetics and oxidative metabolism will impair insulin secretion; for example the mitochondrial Complex I inhibitor rotenone (MacDonald & Fahien, 1990; Zhou et al., 1999), Complex III inhibitor antimycin (Meglsson et al., 1987; MacDonald & Fahien, 1990), F_0F_1 ATPase inhibitor oligomycin (Zhou et al., 1999; Ortsäter et al., 2002) and protonophore uncoupler FCCP (MacDonald & Fahien, 1990; Ortsäter et al., 2002) all act to achieve this.

The ADs, amitriptyline, amoxapine, citalopram, clomipramine and fluoxetine have all been shown to cause mitochondrial dysfunction in other cell systems (Weinbach et al., 1986; Eto et

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3 al., 1985; Souza et al., 1994; Xia et al., 1999; Curti et al., 1999; Chan et al., 2005). Where such
4 events can culminate in oxidative stress and induction of apoptosis (Abdel-Razaq et al., 2011; Li
5 et al., 2012).

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8 Mitochondria have key roles in the function of many other cell types, in particular those with a
9 high energetic demand, for example sensory cells, neurons, skeletal and cardiac muscle. Why
10 reports of adverse effects of chronic AD usage only appear for pancreatic β -cell function and not
11 that of other tissue suggests that the mitochondria of the β -cell may be unique in its sensitivity to
12 these classes of drugs. Indeed the predicted side effects of mitochondrial dysfunction in other
13 cell types, whose function depends heavily on this organelle function: impaired vision, hearing
14 loss, cardiac failure and skeletal muscle weakness, are not normally associated with AD usage
15 (Joint Formulary Committee., 2016).

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18 Although mitochondria have key roles in many cell types, whether the diabetogenic effect of
19 ADs is indeed mediated by action of these drugs on this organelle in pancreatic beta-cells has not
20 been thoroughly explored. In this study, the ability of three ADs: paroxetine, fluoxetine and
21 clomipramine, with particular emphasis on fluoxetine, to disrupt mitochondrial function, impair
22 bioenergetics and inhibit insulin secretion of native and cell-line models of murine pancreatic β -
23 cells was investigated. In addition we explored both the effect of chronic actions of these drugs
24 in order to mimic their long term usage, as well as their acute actions at higher concentrations in
25 an attempt to mimic overdose and drug-drug interaction situations.

36 37 38 39 **Material and Methods**

40 41 **Cells**

42 Unless stated otherwise, cells of the mouse pancreatic β -cell line MIN6 were employed
43 (Miyazaki et al 1990). The MIN6 cell line is a well-established, time honored, model for the
44 study of pancreatic β -cell function (Miyazaki et al., 1990; Ishihara et al., 1993; Daunt et al.,
45 2006; Cataldo et al., 2015). The main advantage of a cell line for pancreatic β -cell research is
46 that it large numbers of phenotypically homogenous cells are available which are devoid of
47 paracrine and neuroendocrine effects that arise from interaction with the other endocrine and
48 neuronal cell types that reside within the pancreatic islet (Ishihara et al., 1993). Although MIN6
49 have been reported to contain and secrete other pancreatic endocrine hormones, these cells still
50 remain primarily β -cell in function (Nakashima et al., 2009). Cells were grown, and the chronic
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3 experiments performed, in Dulbecco's modified Eagle's medium (DMEM) which contained 25
4 mM glucose and was supplemented with: 2 mM L-glutamine, 10% fetal calf serum, 50 μ M 2-
5 mercaptoethanol and 25 mM HEPES. In this study cells from passage numbers 35-42 were used.
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7 Some experiments were repeated with the human pancreatic beta-cell line, 1.1B4 (McCluskey et
8 al., 2011). These cells were maintained and handled using the same protocols as for MIN6
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10 For validation in native tissue we used β -cells both from and within mouse islets of Langerhans.
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12 Pancreatic islets were isolated from the pancreata of male 30-35g CD1 mice as previously
13 described (Daunt et al., 2006). Animals were kept under conditions and methods used that
14 complied with local University rules approved by the UK government home office.
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20 21 **Chronic studies**

22 **Measurement of cell viability with MTT**

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25 The MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide) assay relies on
26 cellular reduction of the MTT substrate to produce a blue formazan product. As such it is a facile
27 indicator of cell viability that results from the product of cell number and redox status (Elmorsy
28 et al., 2014). The MTT assay was employed in accord with the manufacturer's protocol
29 (CellTiter 96 Non-Radioactive Cell Proliferation Assay; Promega, UK). Cells were incubated for
30 4, 24, 48 or 72 hours in the presence of ADs or vehicle. In one set of experiments, this assay was
31 repeated in the presence of 10 mM reduced glutathione (GSH) with a concentration of AD that
32 inhibited MTT reduction by \sim 50% after 24 hours incubation (Elmorsy et al., 2014).
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40 To account for inter-experimental variation between well cell densities, MTT absorbance values
41 are expressed as a percent of that measured for its respective vehicle control (defined as 100%).
42 For all chronic assays, each experiment was performed in triplicate (technical repeat) with the
43 average value considered as a single biological repeat.
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48 **Intracellular ATP content**

49 As a direct measure of bioenergetic potential, intracellular ATP was measured as previously
50 described (Elmorsy and Smith, 2015). Data was corrected for background luminescence by
51 subtraction of the values measured in media alone. ATP levels are represented as a percentage
52 measured in their respective vehicle control.
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Insulin secretion

For this and all subsequent chronic AD incubation assays, cells at confluence were washed twice with PBS and incubated for 24 hrs in DMEM with ADs or vehicle. To maximize effect detection, concentrations of ADs were chosen that inhibited MTT reduction/ATP production by ~ 50% after 24 hours incubation.

To measure insulin secretion, the media was removed and the cells washed with PBS. Cells were then incubated for 15 minutes in serum free DMEM (25 mM glucose) absent of AD. This was removed for insulin assay and the cells measured for total insulin content. Insulin was measured by ELISA (Crystal Chem, Downers Grove, IL, USA) with secretion normalized to total insulin content.

Lactate production

To measure lactate in the chronic experiments the supernatant was collected and assayed by kit in accord with the manufacturer's instructions (Biovision, Mountain View, CA). Lactate production was normalized to cell number, estimated with a hemocytometer, and expressed as a percentage of lactate produced with the vehicle control.

Mitochondrial complex activity

The effect of ADs on MIN6 cells mitochondrial complexes I and III were assessed as previously described (Elmorsy and Smith, 2015). To maximize event detection, concentrations of ADs were chosen that inhibited MTT reduction/ATP production by ~ 50% after 24 hours incubation.

Oxidative stress

The 3,7-dichlorodihydrofluorescein diacetate (DCFDA) assay was used to detect reactive oxygen species (ROS) and was performed following the protocol of Elmorsy et al (2014). For a positive control 1 μ M Antimycin A was added for 30 minutes, this blocks Complex III and generates of ROS. Non-stained cells were used for the blank and were subtracted from the data to correct for background luminescence.

Caspase activity

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3 To determine if the ADs, may induce apoptosis we evaluated the activities of caspases 3, 8 and 9
4 after treatment with the ADs for 24 hrs. These were determined using a BD ApoAlert caspase
5 fluorescent assay kits (Clontech Laboratories, Palo Alto, CA).
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10 **Acute studies**

11 Unless stated otherwise, acute experiments were performed in a HEPES-buffered Hanks solution
12 which contained (in mM): 138 NaCl, 4.2 NaHCO₃, 1.2 NaH₂PO₄, 5.6 KCl, 1.2 MgCl₂, 2.6
13 CaCl₂, 10 HEPES (pH 7.4 with NaOH) and 0.01 % fatty acid free BSA.
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19 **Mitochondrial respiratory rate: O₂ consumption**

20 The rate of O₂ consumption (OCR) of MIN6 and 1.1B4 cell suspensions were measured
21 polarographically with a redox over potential of -0.625 V using Clark Oxygen Electrodes (Rank
22 Brothers, Bottisham, UK) as described previously (Daunt et al., 2006). The background O₂
23 consumption by the electrode was corrected for by subtraction of the OCR measured after the
24 addition of 6 mM Na azide which blocks Complex IV and mitochondrial respiration. Results
25 obtained with the N,N,N',N'-tetramethyl-p-phenylenediamine (TMPD) and ascorbate electron
26 shuttle system (ASB; Duchen et al., 1993; Daunt et al., 2006) were corrected for autoxidation of
27 ascorbate via subtraction of the OCR obtained with ascorbate in the absence of cells: (0.3 nmol
28 min⁻¹ in 1 ml chamber volume). To improve the statistical power for the substrate experiments,
29 data was pooled from both control, and AD experiments, which resulted in different n numbers
30 as indicated.
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42 **Real time lactate production**

43 The rate of lactate production of MIN6 and 1.1B4 cell suspensions was measured simultaneously
44 in conjunction with OCR. The appearance of lactate in the media was measured
45 polarographically with Sarissa Lactate enzyme Electrodes according to the manufacturer's
46 protocol (Sarissa Biomedical Ltd Coventry, UK). The potential of the lactate electrode was set at
47 +500 mV relative to a Ag/AgCl reference electrode (DriRef, WPI). At this potential the electrode
48 oxidises H₂O₂, the product of lactate oxidation by lactate oxidase immobilised on the surface of a
49 carbon fibre electrode; the resultant electrical current is a stoichiometric measure of lactate
50 concentration. Electrode calibration was performed with known concentrations of lactate and
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3 was linear up to $\sim 800 \mu\text{M}$ (Sarissa Biomedical Ltd). The lactate output of the cells was
4 calculated after subtraction of the linear background leak current of the electrode measured in the
5 absence of the acid ($-1.38 \text{ nmol min}^{-1}$ in 4 ml chamber volume). Compounds were confirmed as
6 not to interfere with the electrode by protocol repetition in the absence of cells.
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10 11 12 **Mitochondrial membrane potential**

13 The mitochondrial membrane potential, $\Delta\Psi_{\text{mit}}$, was monitored with Rhodamine-123 (Rh-123)
14 similar to that described previously (Daunt et al., 2006). Only cells that responded to glucose
15 with a decrease in fluorescence and an increase to the protonophore FCCP (carbonyl cyanide-4-
16 (trifluoromethoxy)phenylhydrazone) were chosen for quantitative analysis. Regions of interest
17 (ROI) were drawn around cells, corrected for background fluorescence by subtraction, and the
18 average ROI fluorescence intensity calculated. ROIs were recorded for both single cells and cell
19 clusters within each visual field. Only cells that responded to glucose (hyperpolarization of
20 $\Delta\Psi_{\text{mit}}$) were used for analysis. Image analysis was performed with custom scripts written in
21 Labtalk (ORIGIN Ver, 6.0, OriginLab Corporation, MA USA).
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31 32 **Patch-clamp studies of K_{ATP} channel activity**

33 As an indicator of sub-membrane ATP concentrations K_{ATP} channel activity was measured the
34 with cell-attached patch-clamp technique as previously described (Ashcroft et al. 1988; Smith et
35 al., 2001). Cells were bathed in a high potassium solution, nominally Ca^{2+} free, to
36 electrochemically clamp the cell membrane potential (V_m) to 0 mV. The pipette potential, V_p ,
37 was held at + 70 mV to clamp the trans-patch membrane potential to -70 mV : ($V_m - V_p$). The
38 pipette solution contained (in mM): 140 KCl, CaCl_2 2.6, MgCl_2 1.2, and HEPES 10 (pH 7.4 with
39 NaOH). 100 μM tetraethyl ammonium was added to block voltage-gated K^+ channels.
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46 Patch pipettes, resistances between 2-4 $\text{M}\Omega$, were drawn from GC150TF capillary glass
47 (Harvard Instruments), coated with dental wax (Kerr) and fire polished before use. Currents were
48 measured using an Axopatch 1D patch clamp amplifier (Molecular Devices). The zero-current
49 potential of the pipette was adjusted with the pipette in the bath just before seal establishment.
50 No corrections have been made for liquid junction potentials ($<4 \text{ mV}$). Currents were low pass
51 filtered at 2 kHz (-3db , 8 pole Bessel) and digitized at 10 kHz using PClamp 9.3 (Axon
52 Instruments, Foster City, USA). Single-channel data were analyzed with half-amplitude
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3 threshold techniques as implemented in Clampfit Ver. 10.6 (Axon Instruments, Foster City,
4 USA). Channel activity, NPo, was recorded for 3 minute periods. These experiments were
5 performed at 21-23 °C a temperature at which glucose metabolism (Dawson et al., 1986) and its
6 ability to block K_{ATP} channels activity (Ashcroft et al., 1988; Smith et al; 2001) in beta-cells is
7 maintained, albeit slower than at 37 °C.
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10 11 **Mitochondrial superoxide**

12 To measure mitochondrial ROS and oxidative stress after acute AD incubation the fluorescent
13 dye MitoSOXTM was employed in accord with the manufacturer's instructions (ThermoFisher
14 Scientific, Waltham MA USA). Cells cultured in 6-well plates to confluence were incubated in
15 the dark with 5 µM MitoSOX reagent for 10 min at 37°C. Cells were then washed three times
16 with PBS and incubated in Hanks solution with drugs for 120 min at 37°C. DMSO was used as
17 the vehicle control, 1 µM antimycin A as the positive control and unstained cells for blank. Cells
18 were then washed again with PBS and the dye excited at 510 nm. The emission at 580 nm was
19 captured by fluorescence microscopy. After subtraction of the blank, the mean value of five
20 random images from each well was taken as single sample. Velocity software (Ver. 6.3.1.0,
21 Perkin Elmer Inc.) was used to analyze the fluorescent intensity of individual cells and clusters.
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33 **Statistical analysis**

34 Statistical analysis and curve fitting was performed using Graphpad PRISM version 7.01 (San
35 Diego, California USA). Data distributions were tested for normality with the D'Agostino &
36 Pearson omnibus normality test and the appropriate statistical test used as given in the text. Inter-
37 experimental variation within the assay are either indicated by plots of raw values or the
38 coefficient of variation, CV.
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43 The concentration-response relationships for the actions of ADs were quantified by fitting the
44 assay data with equation 1: $Y = (100 - \text{pedestal}) / (1 + ([AD] / IC_{50})^h)$
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46 Where Y is the % of control (taken as 100%), pedestal is the minimum value for Y, h is the slope
47 index, [AD] is the concentration of antidepressant and IC₅₀ the concentration of drug that
48 produces half-maximal inhibition. All the data generated was confirmed to best fit with a single
49 rather than a two site binding model by the comparison method of Akaike's Information Criteria
50 (Graphpad PRISM).
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3 The relationship between incubation time and IC_{50} was quantified by fitting the data with the
4 empirical equation 2: $IC_{50, t} = (IC_{50, t=0} - \text{plateau}) \times \exp(-K \times t) + \text{plateau}$

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7 Where $IC_{50, t}$ is IC_{50} at time t in hours, $IC_{50, t=0}$ is the IC_{50} at time 0, plateau is the asymptotic value
8 of IC_{50} and K is the rate constant (hours^{-1}). For the majority of the acute fluoxetine studies its
9 $IC_{50, t=0}$ value was used: 30 μM .

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12 Data are given in text as the mean \pm SEM or median with 5 to 95% confidence intervals (C.I.),
13 with n the number of individual determinations as appropriate. Statistical significance is defined
14 as $P < 0.05$. In graphics, $P < 0.05$ is flagged as *, $P < 0.01$ as **, $P < 0.001$ as *** and $P < 0.0001$
15 as ****. Statistical tests are stated in the text and figure legends where appropriate.
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20 21 **Results**

22 **Chronic studies**

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24 The ability of MIN6 to reduce MTT was maintained up to 72 hours, the longest period used (Fig.
25 1A); in control cells there was no change in MTT reduction with time (Pearson $r = 0.79$, -0.71 -
26 95% C.I; $p=0.21$), indicative of little cell replication over this time frame. Inter-experimental
27 variation was small and ranged between 2.2 to 4.7 % ($n = 9$). Figs. 1B-D illustrates that
28 paroxetine, fluoxetine and clomipramine inhibited the ability of MIN6 to reduce MTT in a time
29 and concentration manner. Equation 1 was best fit to the data with IC_{50} and pedestal values given
30 in Figs. 1E and F ($h \sim 0.5$). The maximum amount of block, as indicated by the decrease in
31 pedestal, significantly increased with incubation time for both paroxetine (F test, $p < 0.005$) and
32 clomipramine (F test, $p < 0.0001$) but not fluoxetine (Fig. 1F). Only the IC_{50} value for fluoxetine
33 showed a significant change in magnitude with time of incubation (F test, $p < 0.0001$; Figs. 1E)
34 and was described by equation 2 with an $IC_{50, t=0}$ of 33 μM (12 to 53, 95% C.I.), a plateau of 4.7
35 μM (0 to 9.5, 95% C.I.) and a K of 0.1 hours^{-1} (0 to 2.8, 95% C.I.). Fig. 1G demonstrates that a
36 significant block of MTT reduction occurred with all three ADs at 100 nM, the lowest
37 concentration used at all incubation times (One sample t test relative to vehicle control). Fig. 1H
38 illustrates that co-incubation with 10 mM GSH partially reversed the inhibition of MTT
39 reduction produced by the ADs.
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55 Fig. 2 shows that intracellular ATP production, like MTT reduction, was inhibited in a
56 concentration-dependent manner after 24 hour exposure to AD. Inter-experimental variation
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3 ranged from 1.7% CV for control up to 1.8 % CV in the presence of fluoxetine (n = 9). Equation
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5 1 was readily fit to the data with the same IC_{50} 's given in Fig. 2D. For each AD tested no
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7 significant difference (ANOVA) was found between the IC_{50} for MTT and ATP; data when
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9 combined with the fact that cell density (Fig 3F) did not change suggests that MTT is measuring
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11 redox status and not viability. All three ADs decreased the activity of both complex I and III
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13 (Fig. 2E and D; One sample t test relative to control). Inter-experimental variation as indicated
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15 by the CV ranged from 5% for control up to 18 % for Complex III in the presence of
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17 clomipramine (n = 3).

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19 Since these ADs decreased the activities of Complexes I and III, and also impaired the ability of
20
21 MIN6 cells to reduce MTT and produce ATP, we investigated whether the drugs could affect
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23 glycolysis via measurement of their lactate output as well as generate ROS. Figs. 3A and B
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25 illustrate that paroxetine, fluoxetine and clomipramine all significantly increased lactate output
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27 and ROS when normalized to their respective controls (One sample t-test relative to 100%).
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29 However, a slightly different picture was observed when inter-plate variability was not
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31 controlled for and raw data was used instead (Figs. 3C &D), with significant effects only
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33 observed for fluoxetine when compared to its respective control (ANOVA with Dunnett's).

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35 A prior 24 hour incubation in paroxetine and fluoxetine, at concentrations estimated to produce ~
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37 50% inhibition of MTT reduction/ATP production, significantly decreased the ability of the
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39 MIN6 cells to secrete insulin (Fig. 3E, One sample t-test relative to control: 100%), however this
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41 treatment did not affect cell density (Fig. 3F). Similar conditions significantly increased the
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43 activity of caspases 3, 8 and 9 (Fig.3G)

44 45 46 **Acute studies**

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48 As it is not readily possible to measure cellular redox potential via MTT reduction or cellular
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50 bioenergetics via ATP production over incubation times of minutes, these parameters were
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52 monitored instead by the rate of respiration (OCR) and mitochondrial membrane potential,
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54 $\Delta\psi_{mit}$, respectively (Duchen et al., 1993). Acute actions of these drugs are in keeping with
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56 accidental drug overdose, loading dosage or short term drug-drug interactions.
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3 MIN6 suspensions consumed O₂ (OCR) at a rate commensurate with cell density at 1.3±0.7
4 nmoles 10⁶ cells⁻¹ min⁻¹ (n = 112). Addition of 10 mM glucose produced a ~1.4 fold increase in
5 OCR to 1.7±0.8 nmols 10⁶ cells⁻¹ min⁻¹ (n = 112), subsequent addition of fluoxetine significantly
6 decreased OCR (Figs. 4A-C). This effect was concentration-dependent with an IC₅₀ of 230 μM
7 (139 to 397, 95% C.I.) and h of 1.1 (0.6 to 1.6, 95% C.I.; Fig. 4D solid line). After only 10
8 minutes incubation, 3 μM was the lowest concentration of fluoxetine to have a significant effect
9 on OCR (Kruskal-Wallis, Dunn's multiple comparison test). Similar observations were made in
10 the human beta-cell line 1.1B4 with an IC₅₀ for fluoxetine of 232 μM (53 to 102, 95% C.I.) and h
11 of 0.8 (0.4 to 1.2, 95% C.I.; Fig. 4D solid line). The apparent potency of the SSRI to inhibit OCR
12 in MIN6 was increased in 100 nM FCCP (F-test, P <0.04; Fig. 4C), with an IC₅₀ of 97 μM (81-
13 117, 95% C.I.; data not shown) and h of 1.9 (0.9-2.8, 95% C.I.). Acutely applied paroxetine failed
14 to affect OCR even up to 100 μM, the highest concentration tested (data not shown) whereas
15 clomipramine inhibited OCR with an IC₅₀ of 284 μM (319-372, 95% C.I.) and h of 1.5 (0.87-2,
16 95% C.I.) with 30 μM the lowest concentration that had a significant effect after 10 minute
17 incubation (Kruskal-Wallis, Dunn's multiple comparison test; data not shown).
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32 To confirm that the acute effect of fluoxetine on O₂ consumption was not unique to cell lines, its
33 effect on mouse pancreatic islets was tested. In the absence of exogenous substrate, pancreatic
34 islets consumed O₂ at 1.9±0.5 pmol O₂ islet⁻¹ min⁻¹ (n = 6). The addition of 20 mM glucose
35 significantly increased OCR ~ 1.6 fold to 3.0±0.4 pmol O₂ islet⁻¹ min⁻¹ (p<0.05; ANOVA,
36 Sidaks's multiple comparison test; Fig. 4D); values akin to those previously reported (Daunt et
37 al., 2006). Incubation in 100 μM fluoxetine for 10 minutes significantly inhibited OCR by 54 %
38 to 1.6±0.3 pmol O₂ islet⁻¹ min⁻¹ (p<0.01; ANOVA, Sidaks's multiple comparison test); a
39 magnitude of block larger than that observed for MIN6 cells at the same drug concentration
40 (33%; Unpaired t test with Welch's correction) and suggests an IC₅₀ of ~100 μM for fluoxetine in
41 the intact islet.
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50 To explore candidate target sites for the acute effect of fluoxetine on beta-cell metabolism, a
51 range of substrates which feed in at different entry points of the catabolic pathway for glucose,
52 stimulate OCR and elicit insulin secretion (Daunt et al., 2006) were tested for their susceptibility
53 to inhibition by this SSRI. These were D-glyceraldehyde, methylpyruvate and α-
54 ketoisocaproate (all at 10 mM), and the electron shuttle TMPD/ascorbate (ASB) (Duchen et al.,
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3 2003). Of these, only glyceraldehyde did not stimulate OCR relative to the osmotic control, 10
4 mM 3-O-methyl glucose, (MOG; Fig. 5A; Kruskal-Wallis, Dunn's multiple comparison test).
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6 Within 10 minutes of incubation, 30 μ M fluoxetine inhibited respiration in the presence of all
7 these substrates, except MOG where the block failed to reach significance, an effect regardless of
8 the substrates' ability to stimulate OCR (Fig. 5B; Mann Whitney test relative to respective
9 vehicle control in each substrate).

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11 Although the magnitude of block produced by fluoxetine was unrelated to the change in OCR
12 produced by substrate, it was positively correlated with the magnitude of basal OCR (Spearman
13 r , Fig. 5C). This is exemplified in Fig. 5D, with the greatest inhibition produced by 30 μ M
14 fluoxetine was in cell suspensions with the largest basal OCR (Spearman $r = 0.64$, $p < 0.0001$).
15 This phenomenon was further explored with different glucose concentrations (Fig. 5E and F).
16 Fig. 5E illustrates that a significant decrease in OCR occurred with fluoxetine ($76\% \pm 2.6\%$; $p <$
17 0.001 for all cases, paired Student's t-test); an effect independent of glucose concentration and
18 the OCR stimulated by the sugar (Spearman $r = 0.04$, $p \sim 0.5$). Because the magnitude of the SSRI
19 effect is independent of glucose concentration, when the effect of the drug is plotted as a % of
20 the OCR measured in glucose, the inhibitory effect of the AD appears to be negatively correlated
21 with the concentration of sugar (Fig. 5F; Spearman $r = -0.96$, $p < 0.01$); with the SSRI appearing
22 apparently less efficacious at the larger rates of respiration associated with higher glucose
23 concentrations.

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39 Fig. 6 demonstrates that 10 mM glucose increased lactate production commensurate with an
40 increase in OCR. Subsequent inhibition of OCR with 30 μ M fluoxetine was associated with an
41 increment in lactate output (Fig. 6A, B). The ability of fluoxetine to block OCR and stimulate
42 lactate output was mimicked by 1 μ M rotenone (Fig. 6C, D), with OCR blocked beyond basal in
43 some cases. Fluoxetine at 30 μ M also inhibited OCR and simulated lactate output in the human
44 beta-cell line, 1.1B4 (Fig. 6E, F). These effects were significantly larger than those seen in MIN6
45 cells, a fact due to the greater rate of glucose utilization by 1.14B.

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53 Fig. 7A shows that in the absence of exogenous substrate the mitochondrial membrane potential,
54 $\Delta\Psi_{mit}$, as indicated by Rh123 fluorescence, was stable. Within one minute of 10 mM glucose
55 addition, a sustained decrease in fluorescence occurred; an effect consistent with energization of
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3 the mitochondria and polarization of $\Delta\Psi_{mit}$ associated with enhanced mitochondrial electron
4 transport due to glucose metabolism (Duchen et al., 1993). Application of $2.5 \mu\text{g ml}^{-1}$ oligomycin
5 decreased fluorescence consistent with an increase in $\Delta\Psi_{mit}$ due to the block of H^+ leakage
6 through the $\text{F}_0\text{-F}_1\text{-ATPase}$ (Duchen et al., 1993). Subsequent addition of $1 \mu\text{M}$ antimycin
7 increased Rh123 fluorescence; an observation consistent with the abolition of mitochondrial
8 electron transport and collapse of $\Delta\Psi_{mit}$ since subsequent addition of $1 \mu\text{M}$ of the mitochondrial
9 protonophore FCCP did not elicit a further increase in fluorescence (Fig. 7A). Fig. 7B shows that
10 after energisation of $\Delta\Psi_{mit}$ with glucose, $30 \mu\text{M}$ fluoxetine increased Rh123 fluorescence;
11 however, the depolarization of $\Delta\Psi_{mit}$ produced by fluoxetine was incomplete as demonstrated
12 by the collapse of $\Delta\Psi_{mit}$ that occurred on the addition of $1 \mu\text{M}$ FCCP. Addition of $1 \mu\text{M}$
13 rotenone, like antimycin A, depolarized $\Delta\Psi_{mit}$, and prevented further dissipation of $\Delta\Psi_{mit}$ by
14 $30 \mu\text{M}$ fluoxetine and $1 \mu\text{M}$ FCCP (Fig. 7C). H_2O , the vehicle for fluoxetine, did not affect
15 Rh123 fluorescence, whereas $1 \mu\text{M}$ FCCP could still collapse $\Delta\Psi_{mit}$ (Fig. 7D). Since the
16 absolute value of $\Delta\Psi_{mit}$ reflects a balance between its generation by electron transport and its
17 dissipation, predominantly via $\text{F}_0\text{-F}_1\text{-ATPase}$ activity (Duchen et al., 2003), the ability of an
18 agent to depolarize $\Delta\Psi_{mit}$ can be due to either inhibition of its generation or/and potentiation of
19 its dissipation. On the basis of the retained ability of fluoxetine to inhibit OCR in the presence of
20 FCCP, we assumed the former and quantified the action of fluoxetine on $\Delta\Psi_{mit}$ by its initial rate
21 of effect (ΔRh123 ; Fig. 7E). ΔRh123 was positively associated with drug concentration (Pearson
22 $r = 0.972$, $p < 0.01$), with $10 \mu\text{M}$ the lowest fluoxetine concentration deemed to have a
23 significant effect (ANOVA, Dunnett's post comparison test). Although $30 \mu\text{M}$ fluoxetine
24 depolarized $\Delta\Psi_{mit}$ relative to H_2O , the vehicle control ($p < 0.01$, Kruskal-Wallis Dunn's multiple
25 comparison test), the magnitude of this effect was unrelated to the extent by which glucose
26 energized $\Delta\Psi_{mit}$ (Pearson $p > 0.3$), nor did it reverse the effect of glucose on $\Delta\Psi_{mit}$ (Fig. 7F).
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48 Fig. 8 shows that $1 \mu\text{M}$ Antimycin A after 30 minutes, but not $30 \mu\text{M}$ fluoxetine after 120
49 minutes, significantly increased ROS in MIN6 cells relative to vehicle control (DMSO; $p < 0.001$,
50 ANOVA Holm-Sidak).
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56 **FCCP, but not fluoxetine reactivates K_{ATP} channel activity**
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3 Fig. 9A show that in the absence of glucose, cell-attached patches displayed single-channel
4 currents with biophysical metrics characteristic for K_{ATP} -channels: a single channel current
5 amplitude (i) of 4.8 ± 0.1 pA, burst kinetics and a mean open-channel dwell time of 2.2 ± 0.1 ms
6 ($n = 27-37$; Ashcroft et al. 1988; Smith et al., 2001). Fig. 9A and 9B shows that addition of 10
7 mM glucose significantly reduced NPo by 99% (42 to 0%, 95% C.I; $p < 0.0001$ Wilcoxon Signed
8 rank test, $n = 46$) relative to the perfusion control that was without effect (Wilcoxon Signed rank
9 test, $n = 10$). Fluoxetine (1-100 μ M) failed to reactivate K_{ATP} channel activity blocked by
10 glucose, even after 10 minutes the longest period tested (Figs. 9A, 9C). Although 1 μ M rotenone,
11 a mitochondrial toxin which mimics the metabolic effects of fluoxetine, nor 1 μ M antimycin,
12 failed to reactivate the channel, 100 nM FCCP did (Kruskal-Wallis; Fig 9A and 9C) confirmed
13 by the subsequent abolition of activity with 20 μ M tolbutamide, a potent selective inhibitor of
14 this channel type (Smith et al., 2001).
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24 Discussion

25 *Antidepressants are cytotoxic to MIN6*

26 Since all three ADs decreased MTT reduction and the ability of the β -cells to maintain ATP
27 levels with similar IC_{50} s without affecting cell density suggest they act to impair redox status.
28 Previous studies have demonstrated that fluoxetine (Caminada et al., 2006; Levkovitz et al.,
29 2005; Krishnan et al., 2008), clomipramine (Levkovitz et al., 2005; Abdel-Razaq et al., 2011)
30 and paroxetine (Amit et al. 2009) inhibit cell metabolism with similar IC_{50} values to those we
31 report. Although these IC_{50} 's exceeded the upper therapeutic concentration, in our hands all three
32 drugs produced cytotoxic effects at the therapeutically relevant concentration of 100 nM. Despite
33 intra-islet concentrations for these drugs being unavailable, magnetic resonance spectroscopy of
34 ^{19}F fluoxetine in the human brain reveals a 10-20 fold accumulation of fluoxetine relative to
35 plasma after 6-12 months treatment (Karson et al 1993; Bolo 2000), Together, these findings
36 help explain the diabetogenic risk associated with prolonged AD usage and exposure (Lustman
37 et al., 1997; Rubin et al., 2008; Brown et al., 2008; Derijks et al., 2008; Andersohn et al., 2009;
38 Kivimäki et al., 2010; Khoza et al., 2011).
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51 *The effects of the ADs are not mediated by blockade of serotonin transport*

52 Fluoxetine and paroxetine are SSRIs, whereas clomipramine acts both as an SSRI and a
53 noradrenaline re-uptake inhibitor. These drugs are thought to achieve their antidepressant
54 therapeutic activity, in part, through blockade of the serotonin transporter (SERT) which
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3 enhances serotonin (5-HT) concentrations in the brain to alleviate the symptoms of depression
4 (Blier & El Mansari, 2013). Pancreatic β -cells secrete 5-HT which may exert autocrine effects
5 (Bennet et al., 2015). However, an involvement of SERT for the cytotoxic effects of these ADs
6 can be dismissed, since β -cells do not express the Slc64a gene that encodes for SERT nor do
7 ADs affect the release of 5-HT from them (Cataldo et al., 2015) .
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14 *Antidepressants impair β -cell mitochondrial function*

15 The IC_{50} for the inhibition of oxidative respiration by acute application of fluoxetine on intact β -
16 cells is comparable to that seen in isolated mitochondria from liver ($\sim 300 \mu\text{M}$; Souza et al.,
17 1994) and brain ($\sim 150 \mu\text{M}$; Curti et al., 1999) both for this drug and its metabolite norfluoxetine.
18 Furthermore, Hroudová et al (2012) reports that fluoxetine inhibits Complex 1 and II with IC_{50} 's
19 of 86 and 266 μM respectively. Other mitochondrial targets for fluoxetine are the F_0F_1 -ATPase
20 (Souza et al., 1994; Curti et al., 1999) and the voltage-dependent anion channel, VDAC (Nahon
21 et al., 2005). However, neither of these proteins are likely targets since when the activity of F_0F_1 -
22 ATPase was uncoupled from mitochondrial electron transport by FCCP, the IC_{50} for fluoxetine
23 was not increased; whereas blockade of VDAC by fluoxetine is associated with an enhanced, not
24 impaired, mitochondrial function (Nahon et al., 2005).
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27 The fact that the IC_{50} values obtained from the acute studies exceed the chronic values probably
28 reflect a failure of the ADs to reach equilibrium over the short time periods employed. This is
29 particularly exemplified by fluoxetine whose affinity increased with incubation time with a half-
30 life of 6.9 hrs.
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33 *Role of mitochondrial Complexes*

34 The observations that AD bioenergetic effects were not associated with changes in cell density is
35 consistent with a decrease in cell oxidative respiration capacity; an idea supported by the loss of
36 mitochondrial Complexes seen in other systems: 30% decrease in Complex IV in the β -cell line
37 INS1E with 48 hour exposure to 1 μM fluoxetine (De Long et al., 2014); >20% decrease for all
38 four Complexes in CHO cells after 18 hour incubation with >10 μM clomipramine (Abdel-Razaq
39 et al., 2011). Moreover, at 1 μM the atypical AD bupropion, also associated with an increased
40 risk of diabetes (Andersohn et al., 2009), impairs mitochondrial Complex IV function and insulin
41 secretion in INS1E cells after a 48 hour exposure (Woynillowicz et al., 2012). Complex I and
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3 complex III inhibition is considered the main source of ROS in cells (Ježek et al., 2005; Chen et
4 al., 2003). Such a process was demonstrated in our in our chronic but not acute studies, an effect
5 also observed in INS1E cells when incubated with fluoxetine (De Long et al., 2014). These data
6 suggest that maintained loss of mitochondrial Complex activity and ROS production only occurs
7 with prolonged AD exposure; effects similar to those observed in β -cells with exogenous H_2O_2
8 (Li et al., 2009).
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10 11 12 13 14 15 16 *Fluoxetine blocks basal β -cell lipid metabolism*

17 In the absence of exogenous substrate pancreatic β -cells respire primarily via beta-oxidation of
18 endogenous lipids (Corkey et al., 1989) which involves FADH and Complex II/III. The
19 observation that acute fluoxetine had an effect size that correlated with basal rather than
20 stimulated respiration lends support to Complex II/III as an acute target for this AD; since
21 inhibition of Complex II/III will prevent beta-oxidation of endogenous lipids and OCR. Indeed,
22 fluoxetine partially mimics the antimycin inhibition of Complex II/III on $\Delta\psi_{mit}$. A
23 mitochondrial target for fluoxetine is supported by the increase in lactate production that occurs
24 with this SSRI, a phenomenon like that produced by rotenone due to a failure of the beta-cell to
25 oxidize NADH via Complex I which necessitates NADH recycling via increased lactate
26 generation from pyruvate.
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37 *Antidepressants decrease insulin secretion by MIN6*

38 The decrease in glucose-stimulated insulin secretion may be due to a combination of both
39 oxidative stress and decreased ATP production. Pancreatic β -cells are characterized by a
40 relatively weak expression of superoxide dismutase, catalase and glutathione peroxidase
41 (Grankvist et al., 1996; Lenzen et al., 1996); enzymes that metabolize and neutralize ROS.
42 Consequently, exposure to oxidative as is the case here, may be sufficient to impair glucose-
43 stimulated insulin secretion (Maechler et al., 1999; Sakai et al., 2003; Li et al., 2009). The
44 finding that co-incubation with GSH partially reversed the ability of AD to inhibit MTT
45 reduction is consistent with this notion. The associated suppression of intracellular ATP due to
46 bioenergetic disruption will also act to decrease insulin secretion as it is the major coupling
47 factor for glucose-induced insulin secretion (Ashcroft et al., 1994; Maechler 2002). The ability of
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ADs to impair beta-cell bioenergetics readily explains their ability to induce hyperglycaemia without hyperinsulinaemia in vivo (Yamada et al., 1999; Gomez et al., 2001).

Fluoxetine fails to activate K_{ATP} channels in MIN6

Plasma-membrane K_{ATP} channel activity was blocked by glucose and solely reactivated by FCCP; however, they were not reactivated by fluoxetine, rotenone or antimycin. This is probably because these compounds failed to impair mitochondrial ATP production sufficiently to reactivate K_{ATP} ; unlike FCCP which uncouples respiration and leads to cellular consumption of ATP via reverse mode F_0F_1 -ATPase activity (Rustenbeck et al., 1997). Rotenone and antimycin are well-known to inhibit insulin secretion independently of K_{ATP} channels due to the intimate relationship between mitochondrial metabolism and secretory steps distal to membrane ionic events (MacDonald & Fahien, 1990; Ortsäter et al., 2002), and as we show here fluoxetine, too can also impair mitochondrial respiration acutely which explains its acute hyperglycemic affects in-vitro (Antoine et al., 2004; Cataldo et al., 2015) and in-vivo (Yamada et al., 1999; Gomez et al., 2001).

Suitability of MIN6 as a metabolic model of native pancreatic beta-cells

We found that glucose stimulated MIN6 OCR 1.4 fold that of basal, close to that of 1.6 observed for mouse islets. Furthermore, the MIN6 cell OCR of $1.7 \text{ nmol } 10^6 \text{ cells}^{-1} \text{ min}^{-1}$ in glucose is similar to that of $3 \text{ nmol } O_2 \text{ } 10^6 \text{ cells}^{-1} \text{ min}^{-1}$ of the mouse islets preparations (Daunt et al., 2006). Moreover, the $2.6 \text{ nmol min}^{-1} 10^6 \text{ cell}$ lactate from MIN6 in 10 mM glucose is similar to that for rat islets: $2.0 \text{ nmol min}^{-1} 10^6 \text{ cell}$ (Sener et al., 1976). Furthermore, we estimate a lactate to OCR ratio of 3.6 ± 1 ($n = 8$) for MIN6, a ratio like that reported for rat β -cells (2.4; Sener et al., 1976). Together, these data support MIN6 cells as a suitable metabolic model of native β -cells in which to study mitochondrial function.

Conclusion and clinical implications

We found that paroxetine, clomipramine and fluoxetine were cytotoxic to mouse pancreatic β -cells, an effect dependent on concentration and duration of exposure. Importantly, adverse metabolic effects were seen at chronic AD concentrations of 100 nM; a value less than the median therapeutic concentration for most dosage regimens (Reis et al., 2009). Moreover, as the

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3 plasma concentration for these ADs are well established to be related to dose (Reis et al., 2009),
4 a daily fluoxetine dose of 80 mg can yield steady state plasma concentrations $>2 \mu\text{M}$ (Lundmark
5 et al., 2001), a concentration sufficient to disrupt beta-cell mitochondrial bioenergetics.
6 Obviously, plasma concentrations during loading and drug-drug interactions concentrations may
7 transiently surpass $2 \mu\text{M}$. Why these ADs seem to have adverse clinical affects manifested
8 predominantly by pancreatic β -cell dysfunction probably relates to the poor antioxidant abilities
9 of this cell type, since they were mitigated in part by GSH. Overall, our data suggests that a
10 direct cytotoxic effect of these ADs on pancreatic β -cells may contribute to the diabetogenic
11 potential of antidepressants.
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20 21 Conflict of interest

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23 The authors declare that they have no conflict of interest
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Figure Captions

Fig. 1 Antidepressants inhibit MTT reduction. **A)** Absorbance of MTT reduction measured (arbitrary units, A.U.) in a murine pancreatic β cell-line (MIN6) under control conditions as a function of time. Each point is a triplicate average from a separate experiment ($n = 9$). **B-D)** Concentration-effect relationships for the effect of paroxetine (**B**), fluoxetine (**C**) and clomipramine (**D**) on the reduction of MTT measured after 4 different times of incubation as indicated: \bullet , 4 hrs; \blacksquare , 24 hrs; \blacktriangle , 48 hrs; \blacktriangledown , 72 hrs. Data are mean \pm SEM as a percentage of values measured under control conditions ($n = 9$ in all cases). Solid lines are fits of equation 1 to the data with the parameter values given in **E**) and **F**). **E-F)** Time courses of IC_{50} and pedestal respectively for the inhibition of MTT reduction by the drugs indicated: \bullet , PAR, paroxetine; \blacksquare , FLX, fluoxetine; \blacktriangle , CLM, clomipramine. For **E**) data is given as mean \pm 95% C.I., for **F**) data is mean \pm SEM ($n = 9$). Solid line in **E** is a fit of equation 2 to the FLX data with parameter values given in the text. **G)** Time course for inhibition of MTT reduction by 100 nM AD taken from **B**, **C** and **D**. Data mean \pm SEM ($n = 9$). The effect of all three ADs are positively correlated with time (Pearson, $r > 0.98$ and $P < 0.05$ for all cases). **H)** Effect of ADs in the absence (open bars) and presence (filled bars) of 10 μ M GSH. Cells were incubated for 24 hours in 55 μ M paroxetine (PAR), 29 μ M fluoxetine (FLX) or 44 μ M clomipramine (CLM); concentrations that inhibit MTT reduction by $\sim 50\%$ over this time period. Data mean \pm SEM ($n = 9$). Statistical significance determined by Paired t-test.

Fig. 2 Antidepressants inhibit ATP production and mitochondrial Complex activity. **a-d)** Comparison of the concentration-effect relationships for three ADs on MTT and ATP levels in MIN6 cells after 24 hours of incubation. Effect of paroxetine (**A**), fluoxetine (**B**) and clomipramine (**C**). Data are the mean \pm SEM as a percentage of control values ($n = 9$ in all cases). Solid lines are fits of equation 1 to the data with the IC_{50} s given in **D**). **D)** Comparison of the IC_{50} 's for the inhibition of MTT reduction (\bullet) and ATP production (\blacksquare) by the drugs indicated: PAR, paroxetine; FLX, fluoxetine; CLM, clomipramine. Data given as mean \pm 95% C.I. Data is from 9 experiments in each case. **E-F)** Comparison between the effects of 55 μ M paroxetine (PAR), 29 μ M fluoxetine (FLX) and 44 μ M clomipramine (CLM) on the activity of mitochondrial Complex I activity (**E**) and Complex III activity (**F**) after 24 hr incubation. Data

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3 shown as mean \pm SEM (n = 3).). Statistical significance determined by One sample t-test with a
4 theoretical mean of 100%.
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10 **Fig. 3 Antidepressants inhibit insulin secretion increase lactate output and promote**
11 **reactive species generation.** Cells were incubated for 24 hours in 55 μ M paroxetine (PAR), 30
12 μ M fluoxetine (FLX) or 45 μ M clomipramine (CLM). Data are shown either as raw values or as
13 a percentage of values measured under control conditions: for lactate production (**A & C**
14 respectively) and generation of reactive oxygen species (ROS) (**B & D** respectively). **E**) Insulin
15 secretion stimulated in response to 25 mM glucose for a 15 minutes period post AD treatment. **F**)
16 Cell count after 24 hr incubation at the concentrations of AD given above. Results are expressed
17 as mean \pm SEM (n = 3 for each situation). **G**) Caspase activity after 24 hr incubation at the
18 concentrations of AD given above. Results are expressed as mean \pm SEM (n = 3-9 for each
19 situation). Open bars, caspase 3; light bars, caspase 8, dark bars, caspase 9. Statistical
20 significance for A, B and G was determined by One way ANOVA with Dunnett's multiple
21 comparisons test relative to control for all others by One sample t-test with a theoretical mean of
22 100%.
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35 **Fig. 4. Fluoxetine inhibits oxygen consumption.** **A**) Representative records of oxygen
36 concentration for three different suspensions of MIN6 cells showing stimulation of oxygen
37 consumption (respiration) by 10 mM glucose and the subsequent inhibition by 10, 30 and 100
38 μ M fluoxetine. **B**) Fluoxetine concentration-response relationship for OCR inhibition as a
39 percentage of that measured in 10 mM glucose for MIN6 (●) and human 1.1B4 (○) cell lines.
40 Data are means \pm S.E.M (n= 6 - 29). See text for details of fitted lines. Statistical significance is
41 relative to vehicle control determined via Kruskal-Wallis for MIN6. **C**) Percentage change in
42 OCR relative to that measured in the absence of exogenous substrate (basal) under the conditions
43 indicated: Glu, 10 mM glucose and then 100 μ M fluoxetine, FLX (Wilcoxon signed rank test, n
44 = 17); Glu, 10 mM glucose, then 100 nM FCCP followed by 100 μ M fluoxetine, FFLX
45 (Friedman test; n = 10). **D**) OCR of 6 different pancreatic islet preparations measured under the
46 conditions indicated: Bas, basal in the absence of exogenous substrate; Glu, in 20 mM glucose;
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3 FLX, 20 mM glucose and 100 μ M fluoxetine. Statistical significance determined by 1-way
4 ANOVA with Sidak's multiple comparison test.
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11 **Fig. 5. Substrate independent effects of fluoxetine on respiration.** **A)** Oxygen consumption
12 rate (OCR) stimulated by methyl-O-glucose (MOG, n = 42), D-glucose (GLU, n = 71), D-
13 glyceraldehyde (GLY, n = 34), methylpyruvate (MP, n = 30) and α -ketoisocaproate (KIC, n =
14 33), all at 10 mM, and by tetramethyl-p-phenylenediamine/ascorbate (ASB; n = 45). Data are
15 box and whisker (Tukey). Statistical significance is relative to MOG determined via Kruskal-
16 Wallis with Dunn's multiple comparison test. **B)** Percentage inhibition of OCR by 30 μ M
17 fluoxetine in the presence of the same substrates as in **A** (n = 8-25). Statistical significance is
18 relative to respective water controls via Mann-Whitney. Data are box and whisker (Tukey). **C)**
19 Spearman r values and associated probabilities for the correlation between magnitude of
20 inhibition produced by 30 μ M fluoxetine and basal OCR (open columns) or OCR stimulated by
21 the substrates indicated (closed columns): MOG (3-O-methylglucose, n = 8), GLU (glucose, n =
22 25), GLY (DL-glyceraldehyde, n = 10); MP (methyl-pyruvate, n = 8), KIC (α -ketoisocaproic
23 acid, n = 10) all at 10 mM and ASB (TMPD/ascorbate, n = 10). **D)** Correlation between the
24 decline in OCR produced by 30 μ M fluoxetine (FLX) and basal OCR (n = 24). **E)** Glucose
25 concentration-response relationships for OCR relative to basal in the absence (\bullet) and presence
26 (\circ) of 100 μ M fluoxetine, (n = 5-8). Data are means \pm SEM. Note the similar difference
27 between the two sets of data for each glucose concentration. **F)** Glucose concentration-response
28 relationships for the % inhibition of OCR by 100 μ M fluoxetine. Data are means \pm SEM and are
29 calculated from that shown **E**.
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46 **Fig. 6. Acute fluoxetine stimulates lactate production.** Simultaneously measured changes in
47 oxygen consumption rate (Δ OCR) and lactate production (Δ Lactate) compared to those
48 measured in the absence of exogenous substrate (basal). **A-B)** Effect of 10 mM glucose (Glu),
49 and then 30 μ M fluoxetine (FLX) in the continued presence of the sugar on OCR (**A**) and lactate
50 output (**B**) from the same 5 sets of experiments (matched by symbol). **C-D)** Effects of 10 mM
51 glucose (Glu), and 1 μ M rotenone (Rot) in the continued presence of the sugar on OCR (**C**) and
52 lactate output (**D**) from the same 5 sets of experiments (matched by symbol). Note that rotenone
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3 inhibits OCR beyond that measured in basal to give a negative value. **E-F**) Comparison of the
4 effects of 30 μM fluoxetine in the presence of 10 mM glucose (Glu) on OCR (**E**) and lactate
5 output (**F**) for MIN6 (MIN6F; $n = 5$) and 1.1B4 (11B4; $n = 5$). Statistical significance is via
6 Mann-Whitney.
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11 **Fig. 7. Rh123 fluorescence as a measure of the mitochondrial membrane potential, $\Delta\Psi_{\text{mit}}$.**

12 Effects of 10 mM glucose and drugs as indicated on $\Delta\Psi_{\text{mit}}$, in MIN6 cells. **A-D**) $\Delta\Psi_{\text{mit}}$ is
13 represented by the change in Rh-123 fluorescence relative to that measured in the absence of the
14 sugar (100%). Compounds were applied for the duration of the bars. Records are representative
15 of at least 3 different experiments for each condition and are formed from the average of at least
16 7 ROIs. **A**) Effects of 10 mM glucose, 2.5 $\mu\text{g ml}^{-1}$ oligomycin (Olig), 1 μM antimycin A (AntA)
17 and 1 μM FCCP (FC). **B**) The effects of 10 mM glucose, 30 μM fluoxetine (FLX) followed by 1
18 μM FCCP (FC). **C**) Effects of 10 mM glucose, 1 μM rotenone (Rot) 30 μM fluoxetine (FLX) and
19 1 μM FCCP (FC). **D**) The effect of water then 1 μM FCCP (FC) in the presence of 10 mM
20 glucose. **E**) Concentration-response relationship for the initial rate of Rh123 fluorescence
21 increase (ΔRh123) produced by fluoxetine (\bullet). Data are the means \pm S.E.M. ($n = 3-9$). Statistical
22 significance is via ANOVA with Dunnett's post comparison test relative to water control. **F**) Box
23 and whisker (Tukey) comparison of Rh123 fluorescence relative to basal (100%); GLU, effect of
24 glucose ($n = 184$); FLX, plus 30 μM fluoxetine ($n = 156$); H2O, water control ($n = 28$). Data is
25 from at least 5 independent preparations in each case, where n is the total number of ROI
26 measured for each condition. Statistical significance is via Kruskal-Wallis with Dunn's multiple
27 comparison test.
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42 **Fig. 8. MitoSOX fluorescence as a measure of mitochondrial superoxide generation. A-C)**

43 Representative images of MIN6 cells in DMSO (**A**), 30 μM fluoxetine (**B**) and 1 μM antimycin
44 (**C**) all for 1 hr from the same experimental run. Red fluorescence demonstrates the detection of
45 the superoxide anion by the MitoSOX dye. Scale bar 30 μm , all images at 50% brightness. **D**)
46 Amount of superoxide generated, measured in arbitrary fluorescence units under the conditions
47 indicated: Ctrl, basal conditions ($n = 8$); DMSO, vehicle control for fluoxetine and antimycin (n
48 $= 5$); FLX, 30 μM fluoxetine ($n = 7$); ANTA, 1 μM antimycin ($n = 9$).
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4 **Fig. 9. Cell-attached study of K_{ATP} channel activity** **A)** Representative records of K_{ATP}
5 channel activity measured from the same MIN6 cell under the successive conditions indicated.
6 Dotted line is the zero-current level, openings are downward. **B)** Box and whisker (Tukey)
7 comparison of K_{ATP} channel activity (NPo) normalised to that measured in zero glucose; Ctrl,
8 perfusion control (n = 10), Glucose, 10 mM Glucose (n = 46). **C)** Box and whisker (Tukey)
9 comparison of K_{ATP} channel activity reactivated, as a % of that blocked by 10 mM glucose, with
10 the treatments indicated; H2O, water vehicle control (n = 8); 1F, 1 μ M fluoxetine (n = 7); 3F, 3
11 μ M fluoxetine (n = 5); 10F, 10 μ M fluoxetine (n = 5); 30F, 30 μ M fluoxetine (n = 8); 100F, 100
12 μ M fluoxetine (n = 5); Rot, 1 μ M rotenone (n = 6); FCCP, 100 nM FCCP (n = 16); Tol, 100
13 nM FCCP + 20 μ M tolbutamide (n = 9); Ant, 1 μ M antimycin (n = 4).
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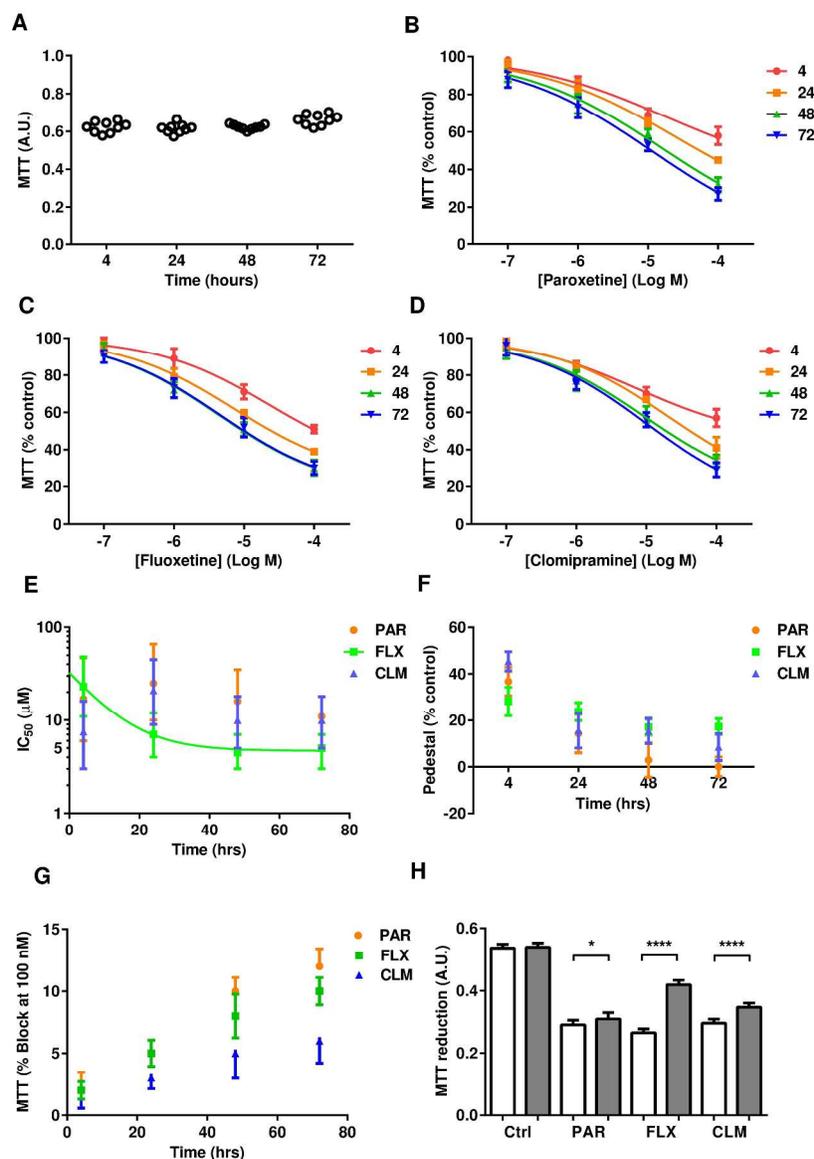


Fig. 1 Antidepressants inhibit MTT reduction. A) Absorbance of MTT reduction measured (arbitrary units, A.U.) in a murine pancreatic β cell-line (MIN6) under control conditions as a function of time. Each point is a triplicate average from a separate experiment ($n = 9$). B-D) Concentration-effect relationships for the effect of paroxetine (B), fluoxetine (C) and clomipramine (D) on the reduction of MTT measured after 4 different times of incubation as indicated: \bullet , 4 hrs; \blacksquare , 24 hrs; \triangle , 48 hrs; ∇ , 72 hrs. Data are mean \pm SEM as a percentage of values measured under control conditions ($n = 9$ in all cases). Solid lines are fits of equation 1 to the data with the parameter values given in E) and F). E-F) Time courses of IC_{50} and pedestal respectively for the inhibition of MTT reduction by the drugs indicated: \bullet , PAR, paroxetine; \blacksquare , FLX, fluoxetine; \triangle , CLM, clomipramine. For E) data is given as mean \pm 95% C.I., for F) data is mean \pm SEM ($n = 9$). Solid line in E is a fit of equation 2 to the FLX data with parameter values given in the text. G) Time course for inhibition of MTT reduction by 100 nM AD taken from B, C and D. Data mean \pm SEM ($n = 9$). The effect of all three ADs are positively correlated with time (Pearson, $r > 0.98$ and $P < 0.05$ for all cases). H) Effect of ADs in the absence (open bars) and presence (filled bars) of 10 μ M GSH. Cells were incubated for

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3 24 hours in 55 μM paroxetine (PAR), 29 μM fluoxetine (FLX) or 44 μM clomipramine (CLM); concentrations
4 that inhibit MTT reduction by $\sim 50\%$ over this time period. Data mean \pm SEM (n = 9). Statistical significance
5 determined by Paired t-test.
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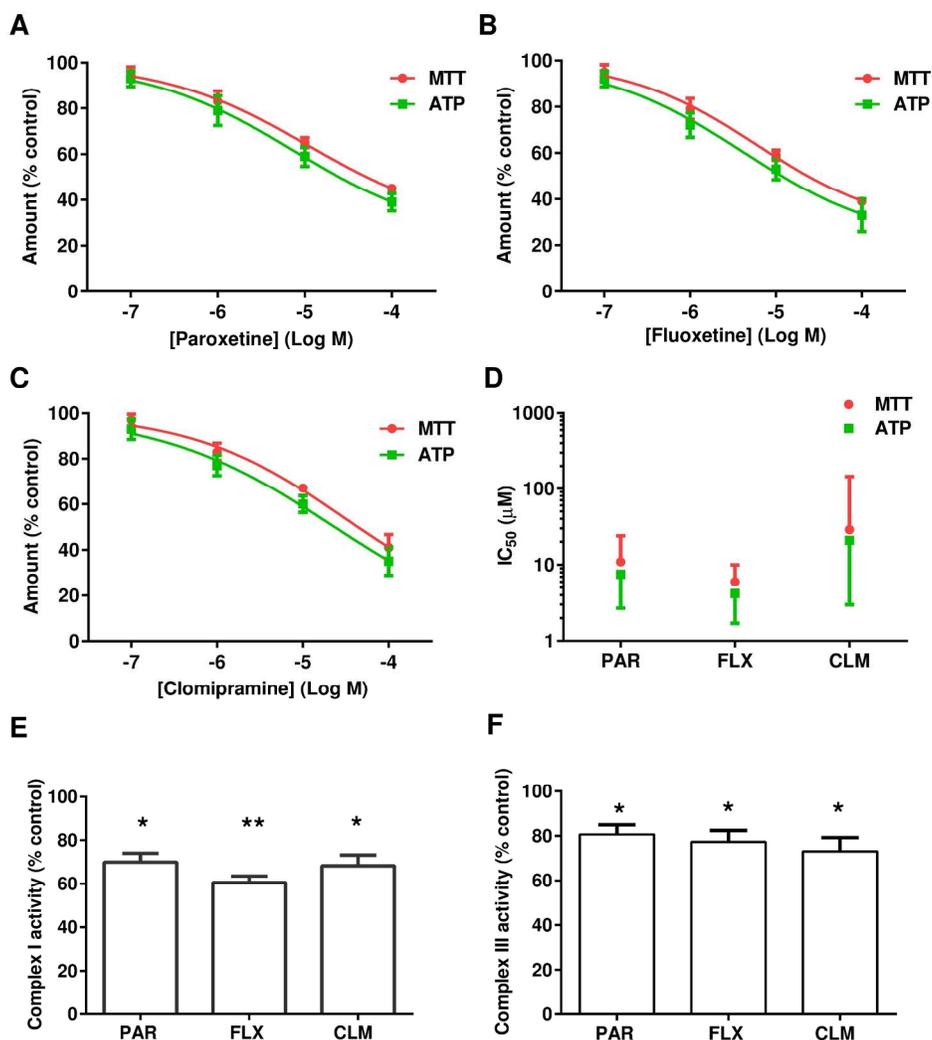


Fig. 2 Antidepressants inhibit ATP production and mitochondrial Complex activity. a-d) Comparison of the concentration-effect relationships for three ADs on MTT and ATP levels in MIN6 cells after 24 hours of incubation. Effect of paroxetine (A), fluoxetine (B) and clomipramine (C). Data are the mean \pm SEM as a percentage of control values ($n = 9$ in all cases). Solid lines are fits of equation 1 to the data with the IC₅₀s given in D). D) Comparison of the IC₅₀'s for the inhibition of MTT reduction (●) and ATP production (■) by the drugs indicated: PAR, paroxetine; FLX, fluoxetine; CLM, clomipramine. Data given as mean \pm 95% C.I. Data is from 9 experiments in each case. E-F) Comparison between the effects of 55 μ M paroxetine (PAR), 29 μ M fluoxetine (FLX) and 44 μ M clomipramine (CLM) on the activity of mitochondrial Complex I activity (E) and Complex III activity (F) after 24 hr incubation. Data shown as mean \pm SEM ($n = 3$). Statistical significance determined by One sample t-test with a theoretical mean of 100%.

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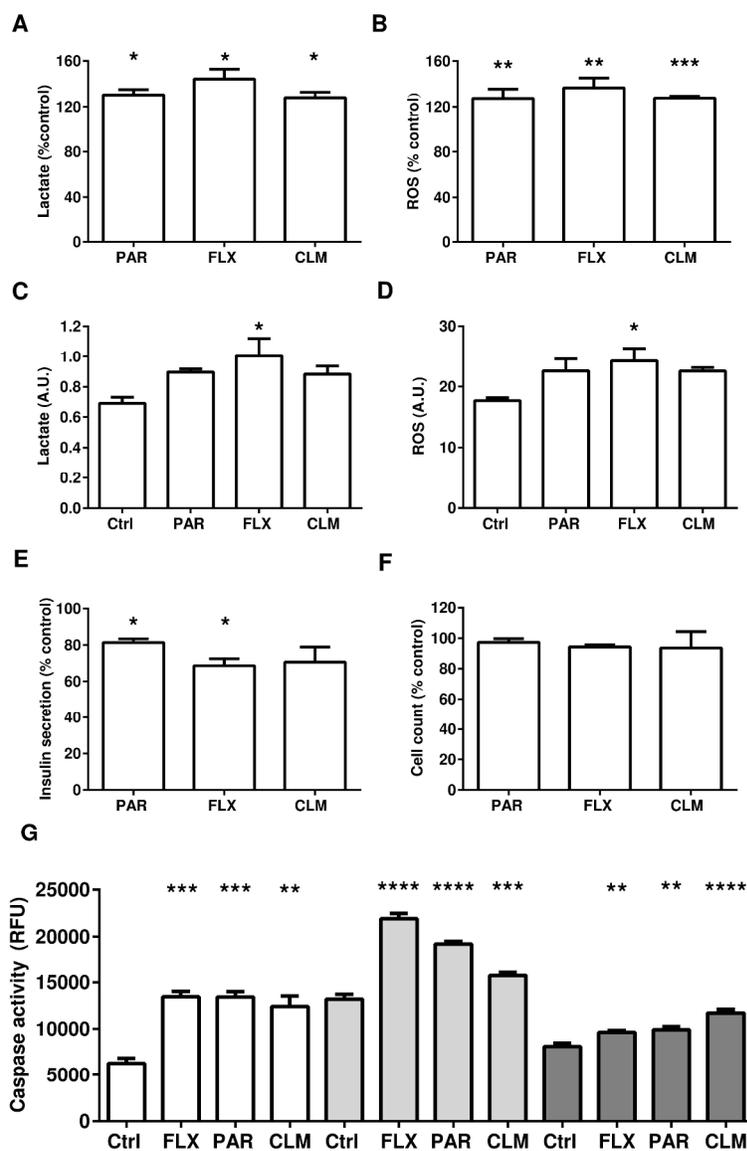


Fig. 3 Antidepressants inhibit insulin secretion increase lactate output and promote reactive species generation. Cells were incubated for 24 hours in 55 μ M paroxetine (PAR), 30 μ M fluoxetine (FLX) or 45 μ M clomipramine (CLM). Data are shown either as raw values or as a percentage of values measured under control conditions: for lactate production (A & C respectively) and generation of reactive oxygen species (ROS) (B & D respectively). E) Insulin secretion stimulated in response to 25 mM glucose for a 15 minutes period post AD treatment. F) Cell count after 24 hr incubation at the concentrations of AD given above. Results are expressed as mean \pm SEM (n = 3 for each situation). G) Caspase activity after 24 hr incubation at the concentrations of AD given above. Results are expressed as mean \pm SEM (n = 3-9 for each situation). Open bars, caspase 3; light bars, caspase 8, dark bars, caspase 9. Statistical significance for A, B and G was determined by One way ANOVA with Dunnett's multiple comparisons test relative to control for all others by One sample t-test with a theoretical mean of 100%.

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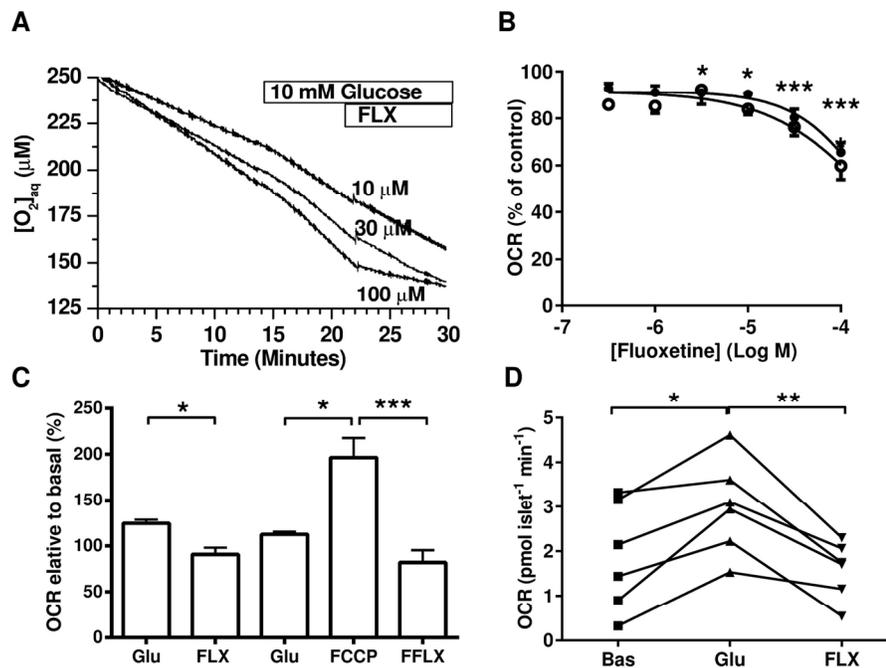


Fig. 4. Fluoxetine inhibits oxygen consumption. A) Representative records of oxygen concentration for three different suspensions of MIN6 cells showing stimulation of oxygen consumption (respiration) by 10 mM glucose and the subsequent inhibition by 10, 30 and 100 μM fluoxetine. B) Fluoxetine concentration-response relationship for OCR inhibition as a percentage of that measured in 10 mM glucose for MIN6 (\bullet) and human 1.1B4 (\circ) cell lines. Data are means \pm S.E.M ($n = 6 - 29$). See text for details of fitted lines. Statistical significance is relative to vehicle control determined via Kruskal-Wallis for MIN6. C) Percentage change in OCR relative to that measured in the absence of exogenous substrate (basal) under the conditions indicated: Glu, 10 mM glucose and then 100 μM fluoxetine, FLX (Wilcoxon signed rank test, $n = 17$); Glu, 10 mM glucose, then 100 nM FCCP followed by 100 μM fluoxetine, FFLX (Friedman test; $n = 10$). D) OCR of 6 different pancreatic islet preparations measured under the conditions indicated: Bas, basal in the absence of exogenous substrate; Glu, in 20 mM glucose; FLX, 20 mM glucose and 100 μM fluoxetine. Statistical significance determined by 1-way ANOVA with Sidak's multiple comparison test.

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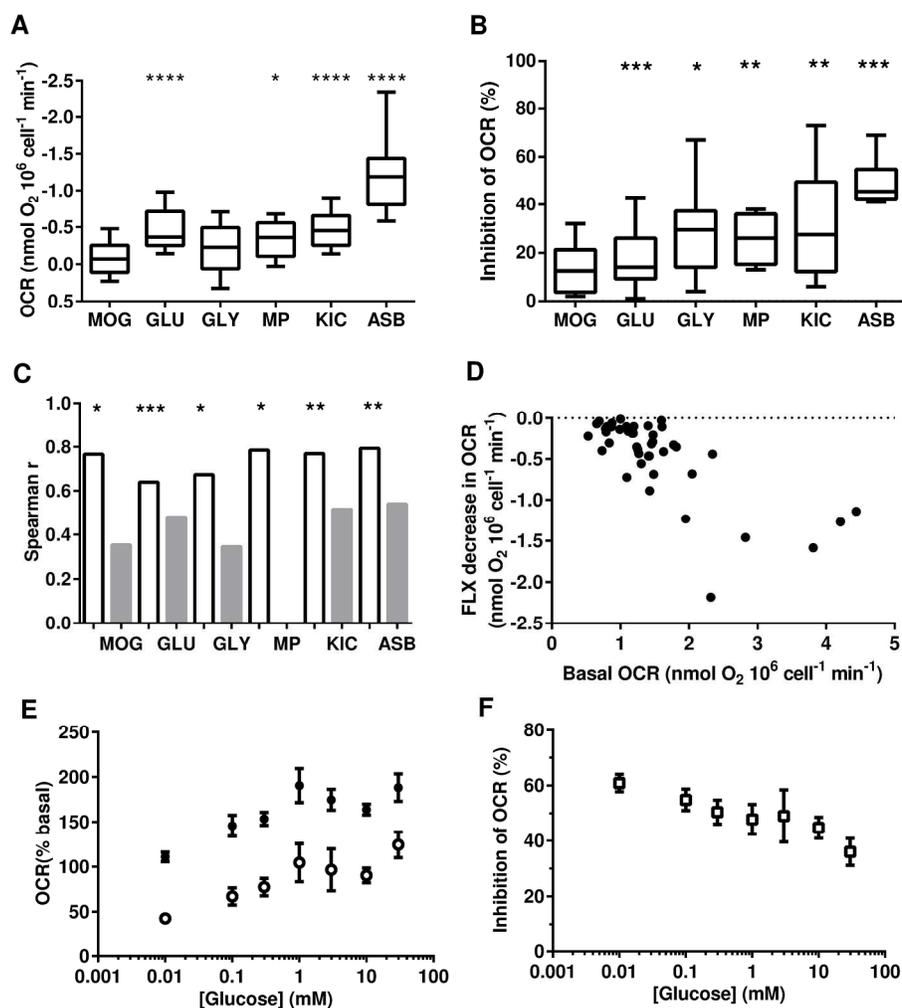


Fig. 5. Substrate independent effects of fluoxetine on respiration. A) Oxygen consumption rate (OCR) stimulated by methyl-O-glucose (MOG, n = 42), D-glucose (GLU, n = 71), D-glyceraldehyde (GLY, n = 34), methylpyruvate (MP, n = 30) and α -ketoisocaproate (KIC, n = 33), all at 10 mM, and by tetramethyl-p-phenylenediamine/ascorbate (ASB; n = 45). Data are box and whisker (Tukey). Statistical significance is relative to MOG determined via Kruskal-Wallis with Dunn's multiple comparison test. B) Percentage inhibition of OCR by 30 μ M fluoxetine in the presence of the same substrates as in A (n = 8-25). Statistical significance is relative to respective water controls via Mann-Whitney. Data are box and whisker (Tukey). C) Spearman r values and associated probabilities for the correlation between magnitude of inhibition produced by 30 μ M fluoxetine and basal OCR (open columns) or OCR stimulated by the substrates indicated (closed columns): MOG (3-O-methylglucose, n = 8), GLU (glucose, n = 25), GLY (DL-glyceraldehyde, n = 10); MP (methyl-pyruvate, n = 8), KIC (α -ketoisocaproic acid, n = 10) all at 10 mM and ASB (TMPD/ascorbate, n = 10). D) Correlation between the decline in OCR produced by 30 μ M fluoxetine (FLX) and basal OCR (n = 24). E) Glucose concentration-response relationships for OCR relative to basal in the absence (\bullet) and presence (\circ) of 100 μ M fluoxetine, (n = 5-8). Data are means \pm SEM. Note the similar difference between the two sets of data for each glucose concentration. F) Glucose concentration-response relationships for the % inhibition of OCR by 100 μ M fluoxetine. Data are means \pm SEM and are calculated from that shown E.

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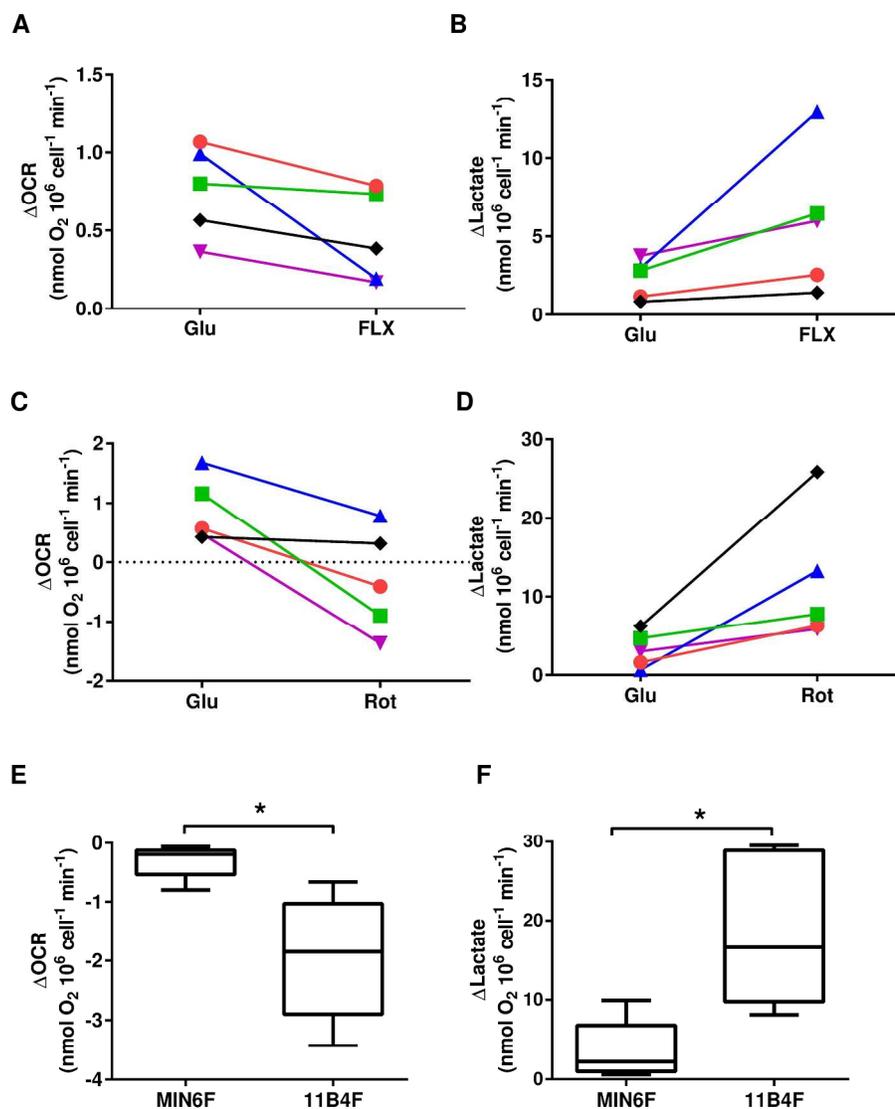


Fig. 6. Acute fluoxetine stimulates lactate production. Simultaneously measured changes in oxygen consumption rate (Δ OCR) and lactate production (Δ Lactate) compared to those measured in the absence of exogenous substrate (basal). A-B Effect of 10 mM glucose (Glu), and then 30 μ M fluoxetine (FLX) in the continued presence of the sugar on OCR (A) and lactate output (B) from the same 5 sets of experiments (matched by symbol). C-D Effects of 10 mM glucose (Glu), and 1 μ M rotenone (Rot) in the continued presence of the sugar on OCR (C) and lactate output (D) from the same 5 sets of experiments (matched by symbol). Note that rotenone inhibits OCR beyond that measured in basal to give a negative value. E-F Comparison of the effects of 30 μ M fluoxetine in the presence of 10 mM glucose (Glu) on OCR (E) and lactate output (F) for MIN6 (MIN6F; $n = 5$) and 1.1B4 (11B4F; $n = 5$). Statistical significance is via Mann-Whitney.

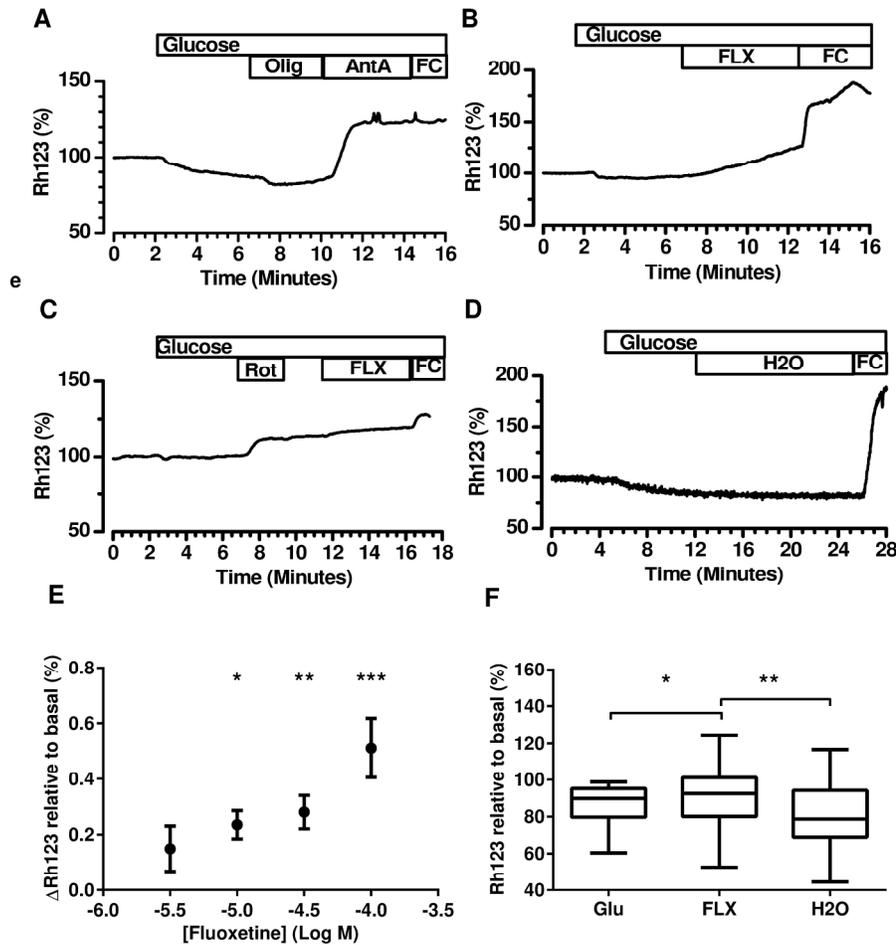


Fig. 7. Rh123 fluorescence as a measure of the mitochondrial membrane potential, $\Delta\Psi_{mit}$. Effects of 10 mM glucose and drugs as indicated on $\Delta\Psi_{mit}$, in MIN6 cells. A-D) $\Delta\Psi_{mit}$ is represented by the change in Rh-123 fluorescence relative to that measured in the absence of the sugar (100%). Compounds were applied for the duration of the bars. Records are representative of at least 3 different experiments for each condition and are formed from the average of at least 7 ROIs. A) Effects of 10 mM glucose, 2.5 μ g ml⁻¹ oligomycin (Olig), 1 μ M antimycin A (AntA) and 1 μ M FCCP (FC). B) The effects of 10 mM glucose, 30 μ M fluoxetine (FLX) followed by 1 μ M FCCP (FC). C) Effects of 10 mM glucose, 1 μ M rotenone (Rot) 30 μ M fluoxetine (FLX) and 1 μ M FCCP (FC). D) The effect of water then 1 μ M FCCP (FC) in the presence of 10 mM glucose. E) Concentration-response relationship for the initial rate of Rh123 fluorescence increase (Δ Rh123) produced by fluoxetine (\bullet). Data are the means \pm S.E.M. (n = 3-9). Statistical significance is via ANOVA with Dunnett's post comparison test relative to water control. F) Box and whisker (Tukey) comparison of Rh123 fluorescence relative to basal (100%); GLU, effect of glucose (n = 184); FLX, plus 30 μ M fluoxetine (n = 156); H2O, water control (n = 28). Data is from at least 5 independent preparations in each case, where n is the total number of ROI measured for each condition. Statistical significance is via Kruskal-Wallis with Dunn's multiple comparison test.

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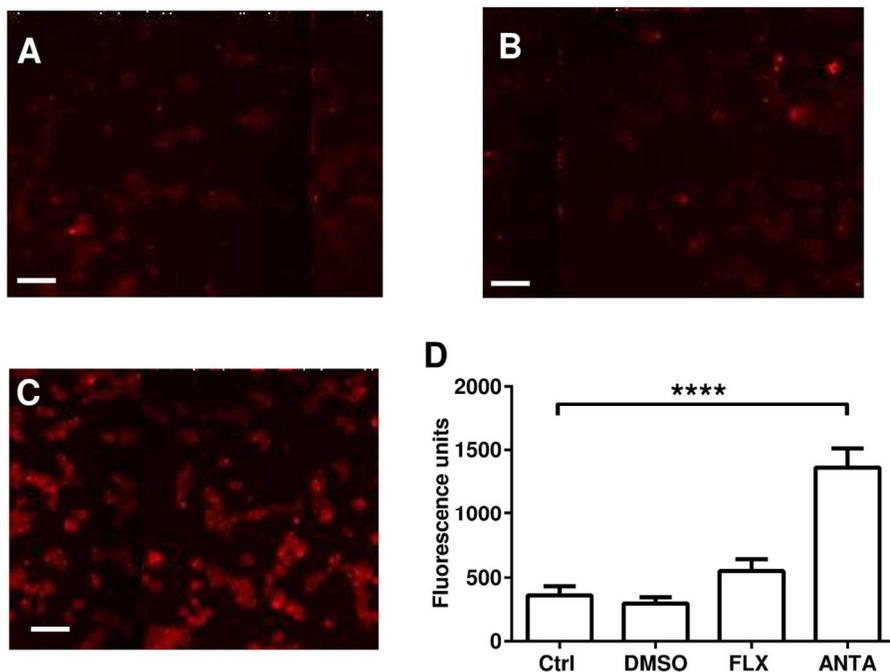


Fig. 8. MitoSOX fluorescence as a measure of mitochondrial superoxide generation. A-C) Representative images of MIN6 cells in DMSO (A), 30 μ M fluoxetine (B) and 1 μ M antimycin (C) all for 1 hr from the same experimental run. Red fluorescence demonstrates the detection of the superoxide anion by the MitoSOX dye. Scale bar 30 μ m, all images at 50% brightness. D) Amount of superoxide generated, measured in arbitrary fluorescence units under the conditions indicated: Ctrl, basal conditions (n = 8); DMSO, vehicle control for fluoxetine and antimycin (n = 5); FLX, 30 μ M fluoxetine (n = 7); ANTA, 1 μ M antimycin (n = 9).

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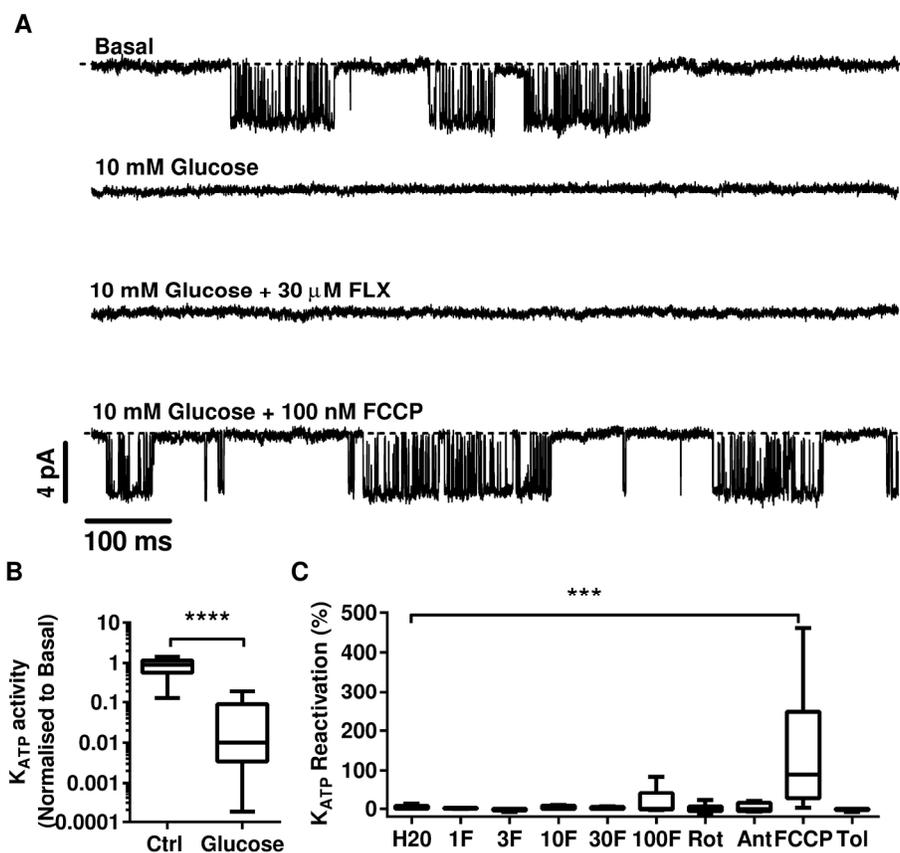


Fig. 9. Cell-attached study of KATP channel activity A) Representative records of KATP channel activity measured from the same MIN6 cell under the successive conditions indicated. Dotted line is the zero-current level, openings are downward. B) Box and whisker (Tukey) comparison of KATP channel activity (N_{Po}) normalised to that measured in zero glucose; Ctrl, perfusion control (n = 10), Glucose, 10 mM Glucose (n = 46). C) Box and whisker (Tukey) comparison of KATP channel activity reactivated, as a % of that blocked by 10 mM glucose, with the treatments indicated; H₂O, water vehicle control (n = 8); 1F, 1 μ M fluoxetine (n = 7); 3F, 3 μ M fluoxetine (n = 5); 10F, 10 μ M fluoxetine (n = 5); 30F, 30 μ M fluoxetine (n = 8); 100F, 100 μ M fluoxetine (n = 5); Rot, 1 μ M rotenone (n = 6); FCCP, 100 nM FCCP (n = 16); Tol, 100 nM FCCP + 20 μ M tolbutamide (n = 9); Ant, 1 μ M antimycin (n = 4).

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