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Original Citation:

Availability:

This version is available at: 11577/3208420 since: 2016-11-16T10:55:09Z

Publisher:

Published version:

DOI: 10.1007/s13300-016-0210-y

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Simvastatin Rapidly and Reversibly Inhibits Insulin Secretion in Intact Single-Islet Cultures

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Received: September 23, 2016

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ABSTRACT

Introduction: Epidemiological studies suggest that statins may promote the development or exacerbation of diabetes, but whether this occurs through inhibition of insulin secretion is unclear. This lack of understanding is partly due to the cellular models used to explore this phenomenon (cell lines or pooled islets), which

are non-physiologic and have limited clinical transferability.

Methods: Here, we study the effect of simvastatin on insulin secretion using single-islet cultures, an optimal compromise between biological observability and physiologic fidelity. We develop and validate a microfluidic device to study single-islet function *ex vivo*, which allows for switching between media of different compositions with a resolution of seconds. In parallel, fluorescence imaging provides real-time analysis of the membrane voltage potential, cytosolic Ca^{2+} dynamics, and insulin release during perfusion under 3 or 11 mM glucose.

Enhanced content The online video files are available to view at <http://www.medengine.com/Redeem/6317F060735F7A3B>.

Electronic supplementary material The online version of this article (doi:[10.1007/s13300-016-0210-y](https://doi.org/10.1007/s13300-016-0210-y)) contains supplementary material, which is available to authorized users.

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Results: We found that simvastatin reversibly inhibits insulin secretion, even in high-glucose. This phenomenon is very rapid (<60 s), occurs without affecting Ca^{2+} concentrations, and is likely unrelated to cholesterol biosynthesis and protein isoprenylation, which occur on a time span of hours.

Conclusions: Our data provide the first real-time live demonstration that a statin inhibits insulin secretion in intact islets and that single islets respond differently from cell lines on a short time scale.

Funding: University of Padova, EASD Foundation.

Keywords: Insulin; Islet; Microfluidic; Simvastatin; Statin

INTRODUCTION

Statins, hydroxy-methyl-glutaryl-CoA (HMG-CoA) reductase inhibitors, are widely used to lower cholesterol and prevent cardiovascular disease [1, 2]. As diabetes is characterized by high cardiovascular risk, international guidelines recommend that most diabetic patients should receive statin therapy [3–5]. Statins effectively reduce cardiovascular morbidity and mortality in patients with diabetes [6], but epidemiological studies suggest they increase the risk of developing diabetes in healthy individuals [7–10] and may worsen glycemia in diabetic patients [11]. Pathophysiologic studies in humans indicate that statins may affect both insulin sensitivity and insulin secretion [12]. In vitro studies show that statins impair insulin secretion by affecting multiple pathways [13], including cellular cholesterol synthesis, membrane fluidity, and isoprenylation of proteins [14]. Though their cardiovascular benefit exceeds the diabetes hazard [15], the mechanisms whereby statins

accelerate the onset of diabetes remain unclear, and whether statins truly exert any action on insulin secretion is debated [16, 17]. This is in part because cellular models used to explore this phenomenon, either cell lines [13, 14] or pooled islets [14], are non-physiologic and lack clinical transferability. Cell lines may behave differently from mature beta cells or have incomplete glucose sensor or insulin secretion machinery, whereas pooled islet cultures are poorly suitable to provide information on dynamic insulin secretion because islet response may be heterogeneous and asynchronous.

A single islet is the minimum fully functional unit of the endocrine pancreas. A precise understanding of islet response to exogenous stimuli is better captured in vitro at the single-islet level, devoid of complex and not fully understood islet–islet interactions. Microfluidics is a technology suitable for cell culture at micrometer scale. Because a single islet has an approximate size of 50–500 μm , it is particularly suitable for microfluidic culture. Accordingly, a number of microfluidic devices have been developed for understanding single islet behavior, as recently reviewed [18]. However the advantages of microfluidic technology go well beyond the size of these devices [19]. For example, Benninger et al. used a microfluidic device to generate precise glucose concentration gradients in a single islet microenvironment to study cell–cell gap junctions in the regulation of coordinated insulin release [20]. Microfluidic devices favor the accumulation of endogenously secreted factors and were previously used to detect islet amyloid polypeptide secretion from as few as ten islets [21] and glucose flux in live myoblasts [22], besides insulin [23].

Here, we study the effects of simvastatin on insulin secretion from single murine islets. A microfluidic system able to temporally control the single-islet soluble microenvironment is

presented. The setup has been technically and biologically validated, before analyzing simvastatin action on intact isolated islets.

METHODS

Microfluidic Setup

The setup was produced by standard photo- and soft-lithographic techniques. Details on the fabrication of the islet culture and the multi-inlet chips are given in the Electronic Supplementary Material (ESM).

Animals and Pancreatic Islet Extraction and Preparation

The protocol followed the Guide for the Care and Use of Laboratory Animals, eighth edition (2011) (<http://grants.nih.gov/grants/olaw/guide-for-the-care-and-use-of-laboratory-animals.pdf>), and was compliant with local and national laws. For murine islet isolation, we optimized a previously described protocol [24]. Details are given in the ESM.

Microfluidic System Running Conditions

The multi-inlet microfluidic system was connected to medium vials of different compositions. Details are given in ESM. This system allowed switching between up to nine different medium compositions by remotely controlling valve opening and closing. The complete system is displayed in Fig. 1.

Islet Staining for Imaging of Viability, Ca^{2+} , Membrane Potential, and Zn^{2+}

Islets were loaded with 1 mg/ml Hoechst 33342 (Sigma-Aldrich) for 10 min. PE-conjugated

Annexin-V (BD Biosciences) was added in the perfusion medium. Prior to microfluidic device insertion, islets were loaded with 3- μM Fura Red acetoxymethyl (AM) (ThermoFisher) or 10 μM Fluo-4 AM (ThermoFisher) for 50 min in RPMI medium at 37 °C for intracellular Ca^{2+} detection; 2- μM FluoZin-3 (ThermoFisher) was added at the same concentration to the different media used for islet stimulation. Extracellular Zn^{2+} concentration detection was used as an indirect measure for insulin secretion detection, as previously described [21]. To avoid biases due to slight differences in the FluoZin-3 concentration or media autofluorescence, before islet insertion into the microfluidic device, a background profile using different media was established at the beginning of each acquisition and then subtracted to the islet response profile. For voltage imaging, islets were incubated for 20 min at 37 °C in RPMI with Vf2.1.Cl (200 nM, kindly provided by Roger Y. Tsien from University of California at San Diego) [25] and Pluronic F-127 (0.1% w/v) (Sigma-Aldrich).

Confocal Imaging

Fluorophores were excited by light from a 460-nm LED (M470L2, Thorlabs) passing through a T510LPXR filter (Olympus) and directed onto the sample through a full mirror (Chroma) while 512 fluorescence emission was selected by a BA495-540HQ filter (Chroma). All fluorescence images were acquired with a 20 \times water immersion objective (NA 0.96, Olympus) and projected on a Complementary Metal-Oxide Semiconductor (CMOS) cooled camera (PCO), controlled by in-house developed software. Image sequences of fluorescence were acquired continuously at three frames per second with 100 ms exposure time.

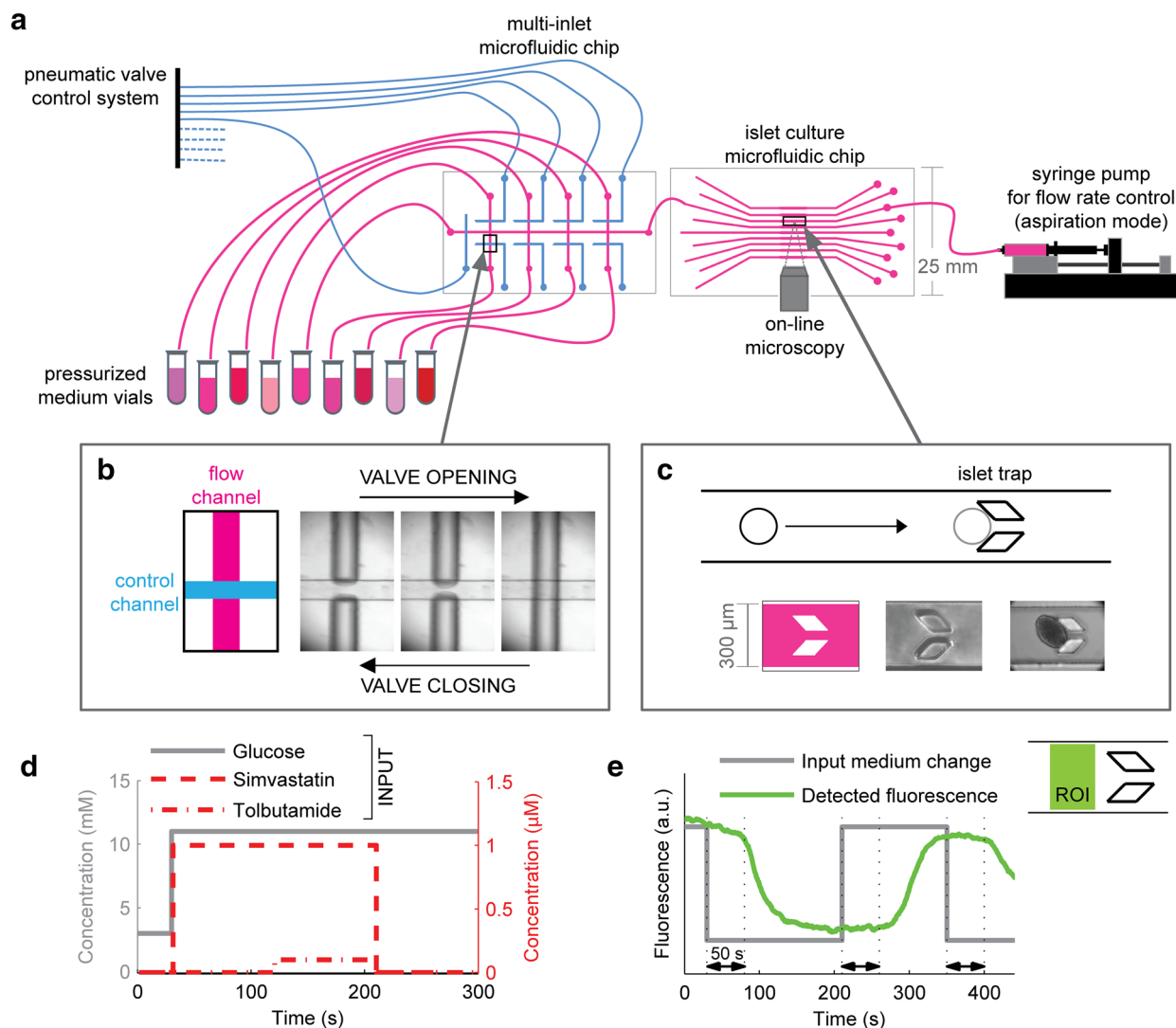


Fig. 1 Microfluidic setup design and validation. **a** Full setup for single islet microscopy observation under controlled dynamic biochemical stimulation. A multi-inlet microfluidic chip contains a system of pneumatic valves that are used to temporally control the medium composition entering the islet culture chamber. The pneumatic valve system is actuated by means of electrovalves (not shown) and is responsible for each valve opening and closing according to the mechanism shown in **c**: pressurizing the control channel prevents medium flow through the flow channel. A single islet is placed in each channel of the islet culture microfluidic chip, where it is precisely positioned because of the presence of a mechanical trap that stops its flow along the channel, as shown in **c**. The islet culture microfluidic chip is a transparent device that is

put under the microscope for online observation. Once the specific valve is open, medium flows through the channel because it is contained in a pressurized vial; to accurately control its flow rate, a syringe pump working in aspiration is placed downstream of the culture system. **d** The system described in **a–c** makes possible the biochemical stimulation of a single-islet with temporal accuracy of seconds. Medium composition is changed over time with minimal external perturbation. **e** Validation of the dynamic input system using medium with and without fluorescein. The theoretical curve (*red*) is compared with the actual medium fluorescence (*green*) detected in the region of interest (ROI) indicated. A delay of approximately 50 s was obtained because of the distance between the valve system and the ROI

Two-Photon Imaging

Two-photon acquisitions were performed with a modular multiphoton microscope (Bergamo II, Thorlabs) equipped with an 8-kHz resonant scanner, extended-field-of-view collection optics, four independent detection channels in the backward direction, and laser-scanned Dodt contrast in the forward direction. The microscope was coupled to two synchronized pulsed laser beams generated by a Ti:sapphire pump laser (Chameleon Ultra 2, Coherent) and an optical parametric oscillator (Camelion Compact OPO). Two-photon microscopy at 800 nm excitation was used to visualize the intracellular calcium (Fluo-4, green), nuclei (Hoechst 33342, blue), and islet structure simultaneously (transmitted light, gray).

Data Analysis

The analysis software, developed in-house in Matlab environment (Release 14, The MathWorks, Inc., Natick, MA, USA), was designed to plot averaged time responses from specific regions of interest (ROIs) as well as to create false-color animations reporting time-dependent fluorescence intensity variations. False-color images were generated using the hue-saturation-value (HSV) visualization algorithm: hue was used to represent fluorescence changes; saturation was set to 1.0; the value (brightness) carried pixel intensity from a reference image that was either updated on a frame-by-frame basis or obtained as an average over a specified number of frames. The resulting frames so constructed were converted to ordinary RGB images and exported in the Audio Video Interlaced (AVI) proprietary video format of Microsoft Video for Windows applications [26]. FluoZin-3 and Fura Red signals were analyzed independently by

playing back images in single-channel mode. ROIs used for Fluo-4 and Fura Red signal analysis were drawn covering the islets, while ROIs for FluoZin-3 signal analysis were drawn outside the islet areas. When Fura Red is excited at 488 nm, its fluorescence emission decreases as Ca^{2+} concentration increases [27]. For this reason, changes in intracellular Ca^{2+} were expressed as $(F_0/F)-1$ ratios [27]. Fluo-4 and FluoZin-3 signals were measured as relative changes of fluorescence emission intensity: $\Delta F/F_0$, where F_0 is the pre-stimulus basal fluorescence; $\Delta F = F - F_0$, F is fluorescence at time t .

Compliance with Ethics Guidelines

The institutional review board of the Venetian Institute of Molecular Medicine approved the study. The protocol followed the Guide for the Care and Use of Laboratory Animals, eighth edition (2011) (<http://grants.nih.gov/grants/olaw/guide-for-the-care-and-use-of-laboratory-animals.pdf>). All institutional and national guidelines for the care and use of laboratory animals were followed.

RESULTS

Microfluidic Single-Islet Culture Setup

The precise dynamic perturbation of a single-islet environment with the online measurement of its response is prevented in conventional culture systems, but can be robustly achieved using microfluidic devices. To this aim, we designed a microfluidic system where a single islet can be trapped without mechanical damage in a defined position, dynamically stimulated with media of multiple compositions, and contemporarily observed under a microscope (Fig. 1a–c). Each

islet was cultured under constant flow perfusion, a condition that better mimics the highly vascularized environment where islets reside in vivo.

The microfluidic device includes an integrated pneumatic valve system (Fig. 1b) that is remotely controlled and enables switching the medium entering the culture chamber in few seconds, while minimizing other culture perturbations (Fig. 1d). The time required for the new medium to reach the islet position within the channel was estimated using buffer with or without fluorescein. At the 2 μ l/min flow rate used in all the experiments, there was an approximate 50 s lag between the medium valve switch and medium change at the islet position (Fig. 1e).

The size and transparency of the microfluidic device make possible the online microscopic observation of the islet response. Overall, the described setup provides maximal information in terms of the spatial and temporal resolution of the collected data.

Biological Validation of the Microfluidic Device

The canonical glucose-stimulated insulin secretion (GSIS) pathway in the β -cell fraction of pancreatic islets involves some well-established sequential steps: glucose is transported into β -cells through plasma membrane glucose transporters and is metabolized, thereby elevating the intracellular ATP/ADP ratio; this event causes the closure of cell-surface ATP-sensitive K^+ channels, leading to cell membrane depolarization; plasma membrane voltage-dependent Ca^{2+} channels are consequently opened, increasing cytosolic Ca^{2+} concentration and triggering the exocytosis of insulin [28].

To interrogate islet behavior via microscopy, we used multiple fluorescent dyes probing cell viability and specific steps of GSIS pathway. We added to the flowing culture media PE-labeled Annexin V, a fluorophore for detecting early apoptosis. We also loaded the islets, prior to their insertion into the device, with dyes enabling the detection of the intracellular Ca^{2+} concentration (Fluo-4 or Fura Red) and cell membrane potential (Vf2.1Cl). We used both confocal and two-photon microscopy.

After each islet insertion into the device, we always observed its morphology to verify the integrity of the external membrane (Fig. 2a, b; Online Video 1). Islets that did not show a visibly intact morphology, for example, due to some variability in the extraction process, were discarded. This occurred in less than 5% of the cases and was usually associated with the presence of some dead cells, assessed by annexin V, not functionally responsive, for instance in terms of intracellular Ca^{2+} oscillations under high-glucose stimulation (Fig. 2c, d).

To verify that the microfluidic system did not impair islet functional response, we perfused the islets with high-glucose medium (11 mM), which is stimulatory for insulin secretion. We used islets from mutant mice that express GFP under the insulin promoter in order to identify the β -cells within the islet. In Fig. 3a, we show persistent oscillations of the intracellular Ca^{2+} at 11 mM. As expected, the oscillations were restricted to β -cells and were not an artifact of the culture system or method of analysis. Moreover, β -cells within the same islet showed a properly synchronized response (Fig. 3b).

To verify islet structural integrity also in terms of cell-cell electrical coupling, we then loaded the islets with Vf2.1Cl, a voltage-sensitive fluorescent dye, and

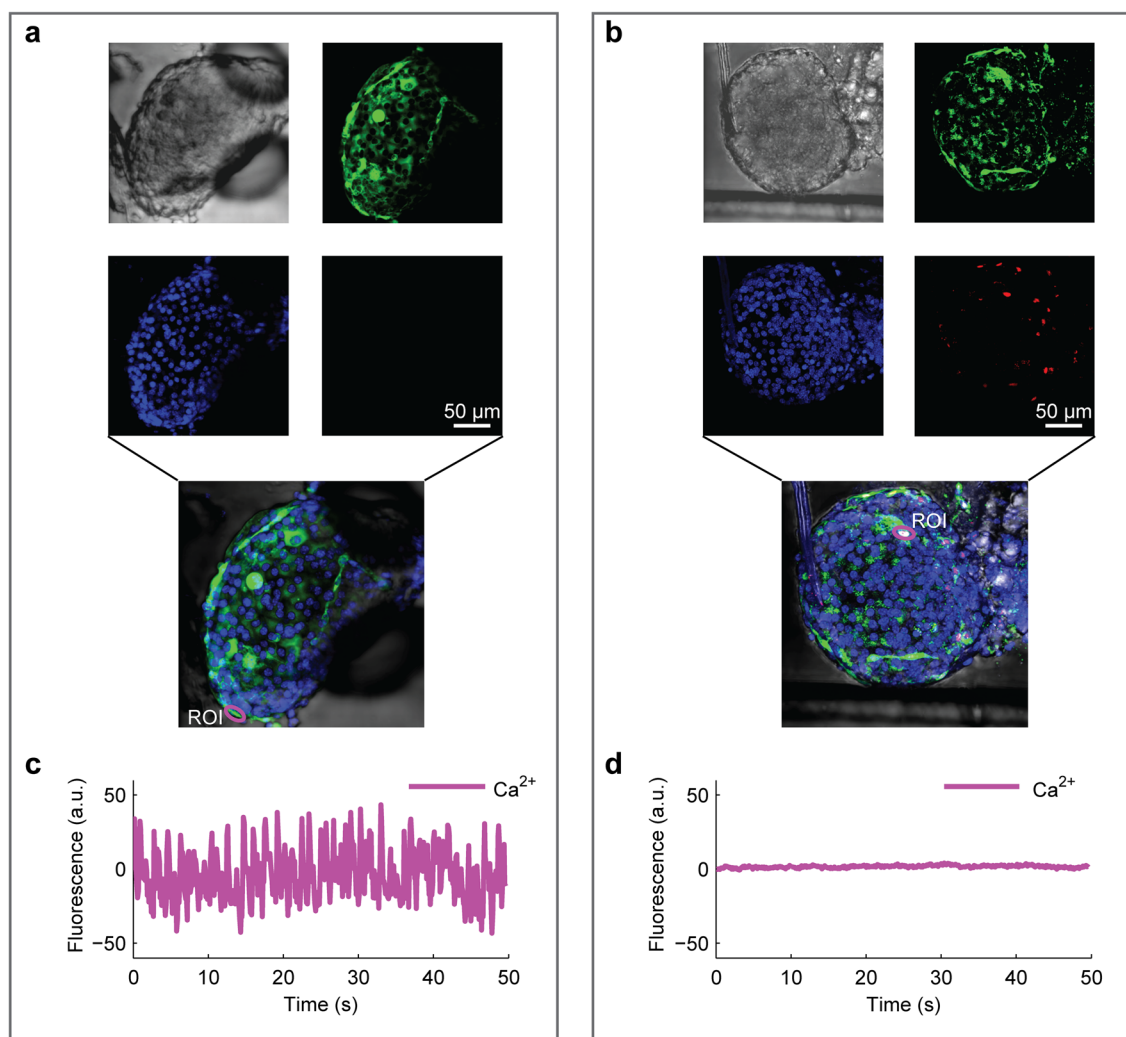


Fig. 2 Two-photon imaging of islet morphology and viability in microfluidics. Islets were pre-loaded with Hoechst 33342 and Fluo-4 for nuclear identification and intracellular Ca^{2+} detection, respectively. Viability was detected by adding annexin V in the flowing medium. **a**, **b** Images of single islets, intact (**a**) or subjected to mechanical stress (**b**), within a microfluidic channel (see video 1). From *top left*: morphology (transmitted light),

performed imaging at high speed. Under high-glucose conditions, we were able to observe the spatial propagation of membrane potential waves throughout the islet (Fig. 3c; Online Video 2), as previously reported [29].

As a last verification of functionality, we loaded the islets with probes for detection of

nuclei (*blue*), intracellular Ca^{2+} (*green*), and mortality (*red*). **c**, **d** Plots of intracellular Ca^{2+} oscillations in the same islets shown in **a**, **b**, respectively, constantly perfused with buffer containing stimulatory glucose concentration (11 mM). Zero-centered raw data. Annexin-V⁺ cells from the stressed islets do not display oscillatory Ca^{2+} activity, as detected by Fluo-4

both membrane potential and intracellular Ca^{2+} . Imaging at high speed, we observed the two waves propagating at the same frequency, but with a small delay of intracellular Ca^{2+} compared to membrane voltage (Fig. 3e, f; Video 3). This result is consistent with Ca^{2+} regulation occurring along the pathway of

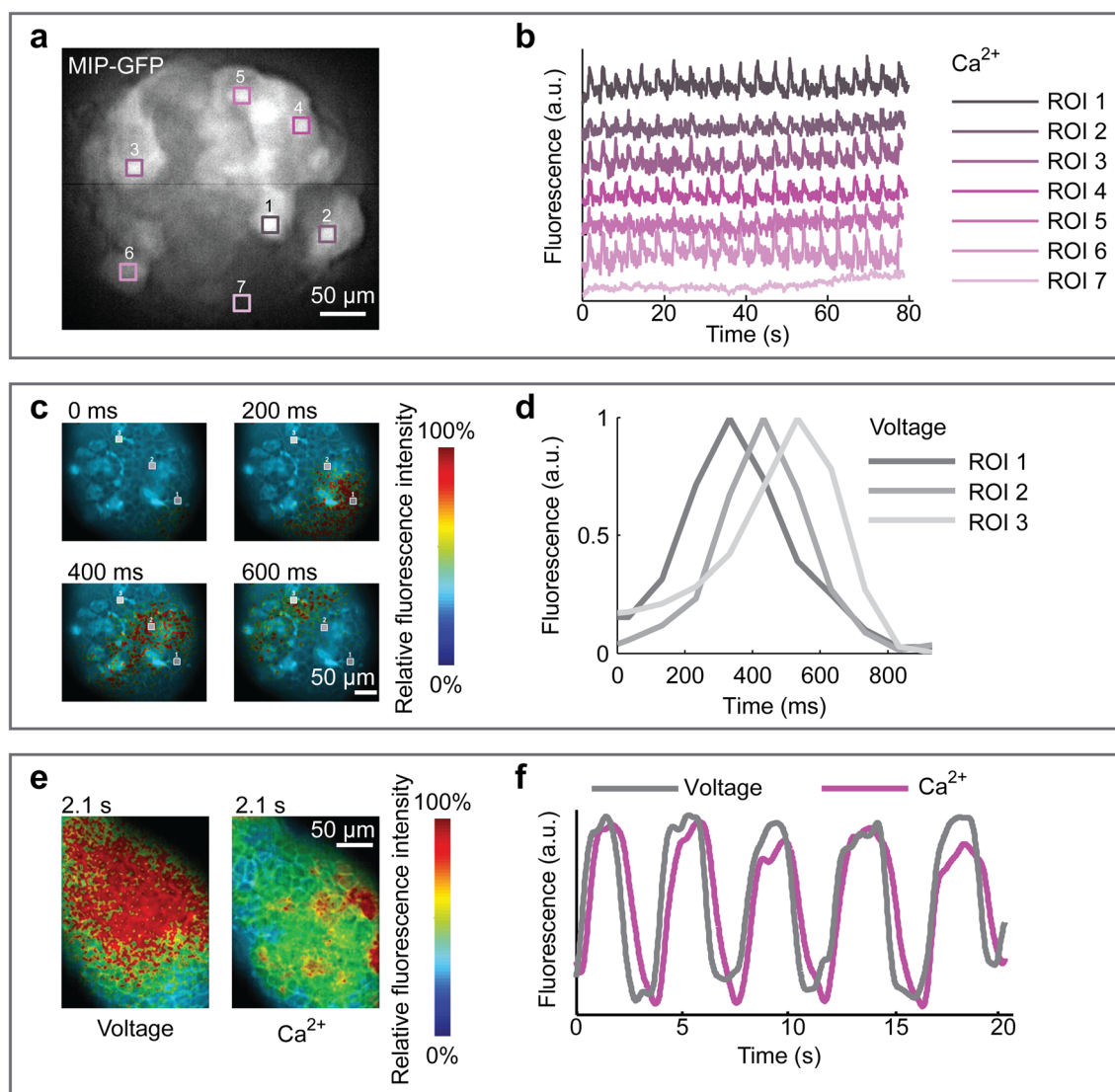


Fig. 3 Validation of islet functional response at 11 mM glucose. **a** Green fluorescence intensity of an islet isolated from a Mouse Insulin Promoter (MIP)-GFP mouse, pre-loaded with Fura Red for intracellular Ca^{2+} detection. **b** Plots of synchronous intracellular Ca^{2+} oscillations of cells in the regions of interest (ROIs) indicated in **a**. ROI 7 refers to non β -cells (GFP), which do not show Ca^{2+} oscillatory activity in response to high glucose concentration. **c** Membrane potential of an islet from a wild-type mouse loaded with the voltage dye Vf2.1.Cl and imaged at the indicated time points during constant perfusion at high glucose. The false-colored image sequence shows

fluorescence intensity. **d** Plot of signal intensity inside ROIs indicated in **c**. Oscillation phase shift describes the spatial propagation of electrical activity from cell to cell within the islet (see also Video 2). **e** Simultaneous recording of intracellular Ca^{2+} and membrane potential in an islet from a wild-type mouse, pre-loaded with Fura Red and Vf2.1.Cl. The false-colored image sequence shows fluorescence intensity. **f** Plot derived from data shown in **e**. Intracellular Ca^{2+} and membrane potential waves propagate with the same periodicity. Ca^{2+} propagation occurs with slightly delayed phase (see also Video 3)

insulin secretion and confirms that each step of the pathway is in place and functional within the microfluidic culture system.

Single-Islet Dynamic Stimulation

Next, we performed dynamic stimulations of the islets, fully taking advantage of the developed microfluidic setup that makes automatically switching between media of different compositions possible. For these experiments, we also added a cell-impermeant Zn^{2+} -sensitive fluorophore (FluoZin-3) to the flowing culture media. Zn^{2+} measurement was used to indirectly evaluate the dynamics of insulin secretion, as Zn^{2+} is present within the intracellular insulin-containing granules undergoing exocytosis [30].

We analyzed single-islet response both upon switching between non-stimulatory (3 mM) and stimulatory (11 mM) glucose concentrations and in the presence of tolbutamide, an ATP-sensitive K^+ -channel blocker that induces insulin secretion even at non-stimulatory glucose concentrations [31]. As expected, tolbutamide produced a sudden increase of insulin secretion (Fig. 4a; Online Video 4) and of intracellular Ca^{2+} concentration (Fig. 4b).

Altogether, these data demonstrate islet functionality within the microfluidic system, validate the detection method for analyzing single islets with high spatio-temporal resolution in response to stimulations, and give a demonstration of the high potential of this system for performing accurate dynamic studies.

Effects of Simvastatin on Insulin Secretion

In a previous study on MIN-6, a murine insulinoma cell line, simvastatin was shown to reduce insulin secretion by targeting

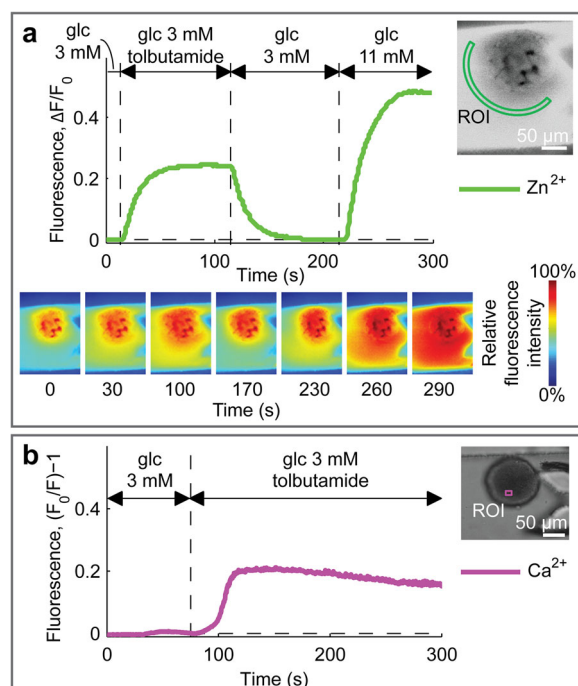


Fig. 4 Physiologic and pharmacologic validation of islet functionality. **a** The islet shown in the inset was dynamically perfused with different media, as indicated, all containing the FluoZin-3 Zn^{2+} -sensitive fluorophore. This dye indicates Zn^{2+} secretion from the islet, as an indirect measurement of insulin secretion [21]. The plot shows changes of fluorescence intensity in the ROI shown in the inset. The false-colored images at the bottom show the spatial fluorescence intensity at different time points (see Video 4). **b** The islet shown in the inset was pre-loaded with FuraRed and dynamically perfused with different media, as indicated. Fluorescence intensity in the inset ROI was quantified in the plot that shows intracellular Ca^{2+} concentration

ATP-sensitive K^+ -channels, voltage-gated Ca^{2+} -channels, muscarinic M3 receptors, and GPR40 in a time span of approximately an hour [13]. On a shorter time scale (minutes), the same study demonstrated that simvastatin suppresses cytosolic Ca^{2+} regulation in response to both stimulatory glucose concentrations and tolbutamide in MIN-6 cells.

We used our validated microfluidic system to analyze short-term effects of simvastatin on the insulin secretion pathway using intact single

pancreatic islets as a more physiologically relevant *in vitro* model.

After testing different concentrations of simvastatin from 1 to 10 μM (data not shown), we used a 1 μM concentration in all experiments, as the lowest concentration of drug that elicited an effect on islet function. We then analyzed the ability of simvastatin to modify insulin secretion by single islets in microfluidics. As shown in Fig. 5a and Online Video 5, simvastatin rapidly blocked insulin secretion even in the presence of a stimulatory glucose concentration (11 mM). The same inhibitory effect on secretion was observed independently of the sequence in which stimuli were applied (Fig. 5a, b). Such a rapid and reversible effect of simvastatin on single islets has never been shown before and was detected taking advantage of the small-scale transparent dynamic setup we developed.

To compare our data to those previously obtained with MIN-6 cells, we coupled the measurement of insulin secretion with that of cytosolic Ca^{2+} . Unlike in Yaluri et al. [13], we found that cytosolic Ca^{2+} was not affected by simvastatin in single islets, despite its inhibition

of insulin secretion at stimulatory glucose concentrations (Fig. 6a; Online Video 6). This suggests that simvastatin acts on the late stages of insulin granule trafficking, downstream of cytosolic Ca^{2+} regulation. However, when tolbutamide was concomitantly used to induce insulin secretion, simvastatin did not prevent a cytosolic Ca^{2+} increase and insulin secretion, at both low (3 mM) and high (11 mM) glucose concentrations (Fig. 6b). Similarly, islet treatment with simvastatin, prior to addition of tolbutamide, did not prevent the rise in cytosolic Ca^{2+} concentration and insulin secretion (Fig. 6c).

DISCUSSION

In this study, we developed and validated a microfluidic platform optimized for functional analysis of single pancreatic islets by live confocal and two-photon imaging. Microfluidic systems are increasingly used to study physiological phenomena *in vitro*, with several advantages [32, 33]. The possibility to analyze a single islet, the fundamental unit of endocrine pancreas, preserves intercellular

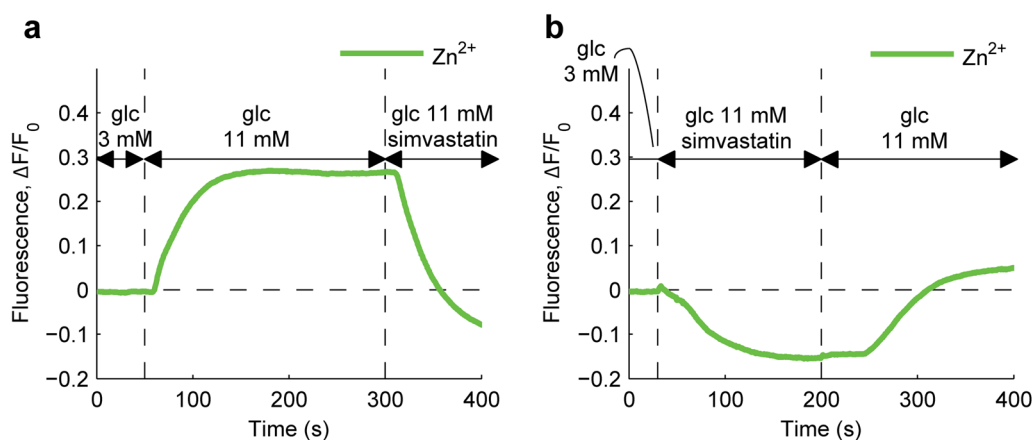


Fig. 5 Effects of simvastatin on glucose-induced insulin secretion. **a** Dynamics of insulin secretion, indirectly measured by FluorZin-3 fluorescence intensity, during islet

perfusion with media of different compositions, as indicated. **b** Same as **a**, inverting the order of simvastatin stimulation at high-glucose concentration

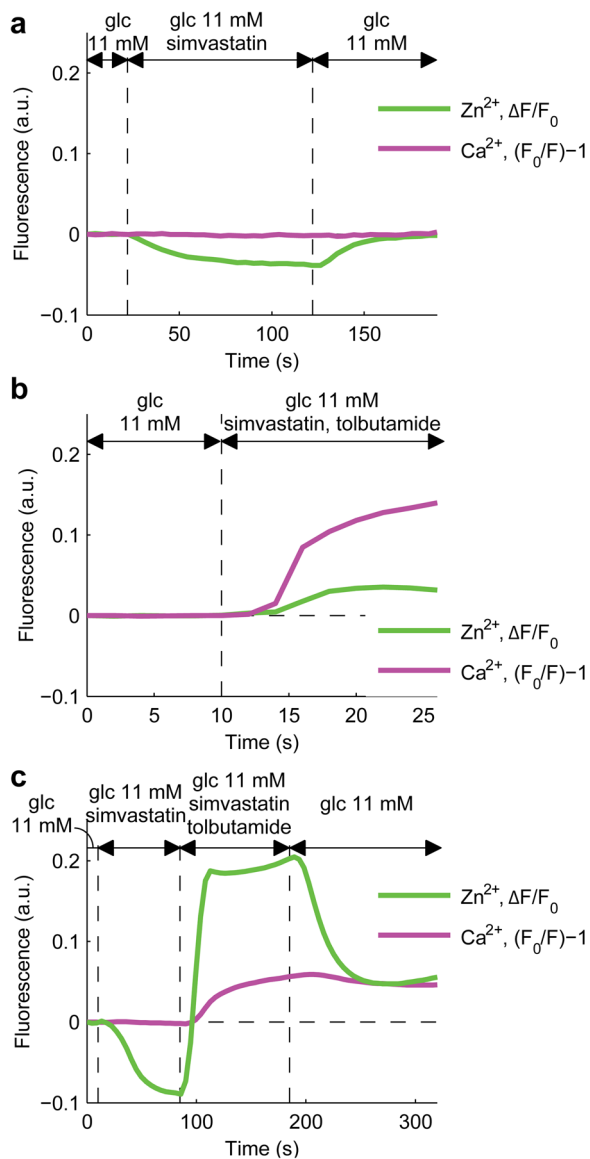


Fig. 6 Effects of simvastatin on insulin secretion and intracellular Ca^{2+} concentration at 11 mM glucose. **a–c** Plots show the simultaneous recording of FluoZin-3 and Fura Red fluorescence intensity of islets pre-loaded with Fura Red and perfused with media of different compositions, as indicated

interactions within the islet and avoids the confounding complexity of whole animal physiology and inter-islet communication as in organo-culture. By using a miniaturized multichannel system we could remotely control the switch of the perfusion medium

entering the microfluidic chamber with a resolution of seconds and analyze live how a single islet responds to different exogenous stimuli. In vivo imaging of the endocrine pancreas is the most suitable physiological model to assess insulin secretion. However, this is poorly standardized, highly invasive, and laborious. In particular, it is difficult to deliver precisely localized stimuli, while dynamically visualizing β -cell response. Ex vivo intact islets are fully functional units, whereas β -cell lines are immortalized and often not pure: the commonly used MIN6 cell line contains a mix of pancreatic endocrine cells [34]. Furthermore, primary islets can be quickly isolated and immediately used for imaging, whereas β -cell lines are difficult to maintain in culture (Table 1).

Using confocal and two-photon imaging, we showed normal functionality of single islets within the microfluidic device in terms of synchronized membrane voltage and intracellular Ca^{2+} regulation, as well as glucose- and tolbutamide-stimulated insulin secretion.

As a clinically relevant application of this setup, we explored the effects of the HMG-CoA reductase inhibitor, simvastatin, on Ca^{2+} dynamics and insulin secretion. Several reports suggest that statins may increase the risk of developing diabetes by acting on insulin secretion, insulin resistance, or both [11, 12, 35, 36]. However, whether statins truly inhibit insulin secretion is unclear.

Pooling together five murine islets, Zúñiga-Hertz et al. demonstrate that simvastatin interferes with isoprenylation of proteins involved in insulin granule exocytosis on a 2-h time scale and with cholesterol biosynthesis on a 24-h time scale [14]. Shorter time effects of simvastatin on the GSIS pathway have only been demonstrated in vitro using a

Table 1 A comparison of different models for beta cell imaging

Methods	Usefulness	Limitation
Cell line	Controlled experimental conditions Reproducibility	Difficult to maintain in culture Immortalized No pure beta cell line (e.g., MIN6) No interaction with alpha/delta cells
Pooled islet culture	Controlled experimental conditions Quick isolation (2 h) Preserved islet heterogeneity integrity High signal-to-noise ratio	Batch-to-batch variability 3 days of vitality after isolation Average response from asynchronous islets
Intact single islet	Controlled experimental conditions Quick isolation (2 h) Preserved islet heterogeneity integrity Quasi-physiological model	3 days of vitality after isolation Islet-to-islet variability
Animal model	High-fidelity physiological model	Difficult visualization No controlled conditions High variability

murine β -cell line, MIN-6, and only as for its action on cytosolic Ca^{2+} dynamics [13].

Here, we extended the results previously obtained with MIN-6 cells to single, freshly isolated murine islets. While Yaluri et al. [13] found that simvastatin affects Ca^{2+} regulation, in our results single islet cytosolic Ca^{2+} is not perturbed by simvastatin. Moreover, for the first time, we were able to demonstrate that simvastatin affects insulin secretion of single islets, a result previously shown only by pooling more islets together, a less physiological model because of the *in vivo* islet–islet coordination [37], and analyzing insulin secretion only on much longer time scales. The rapidity of the simvastatin effect on insulin secretion suggests its involvement in regulatory mechanisms other than cholesterol biosynthesis and

post-translational modifications, which require more time to occur.

Ultimately, we showed that single islets respond to simvastatin acutely and reversibly, blocking the normal insulin secretion induced by high glucose. As there is no effect of simvastatin on Ca^{2+} intracellular concentration and dynamics, we suggest that the inhibitory effect on insulin secretion may be exerted at the levels of insulin granule trafficking. Moreover, tolbutamide can effectively counteract the inhibitory effect of simvastatin on insulin secretion, suggesting that simvastatin's action is reversible not only by its withdrawal, but also by Ca^{2+} overload and membrane depolarization. Reversibility of the effect has clinical implications and likely reflects the relatively benign effect of statins

Table 2 Advancement over state of the art

What was previously known?	Clinical trials and epidemiological studies suggest that statins increase the risk of developing diabetes in healthy individuals and may worsen glycemia in diabetic patients In vitro studies show that statins impair insulin secretion by affecting multiple pathways, including cellular cholesterol synthesis, membrane fluidity, and isoprenylation of proteins
What this study adds?	We have used a new methodological approach to dynamically study the inhibitory effect of simvastatin on insulin secretion of intact islets. Data show this effect is rapid and reversible
What are the implications?	Rapidity of inhibition suggests mechanisms unrelated to cholesterol synthesis, but possibly based on electrophysiology or granule docking/fusion modulation Reversibility of inhibition has clinical implications and may suggest a relatively benign effect of statins on beta cell function
Which are the next steps?	To better dissect the molecular mechanisms of insulin secretion inhibition by statins To identify concomitant treatments that may counteract the effects of statins on insulin secretion

on diabetes onset or progression. Limitations of this study are inherent to the short time course of observation and the lack of a dose-dependent response. Table 2 highlights the advancement over the state of the art made by our study.

CONCLUSION

In conclusion, we report that simvastatin rapidly and reversibly impairs insulin release from intact islets *ex vivo*. Further work is needed to understand the underlying mechanisms of action of simvastatin at the second-to-minute scale. The microfluidic and imaging techniques herein described will be suitable to dissect the molecular and electrophysiological processes underlying this statin effect. The possibility to use concomitant treatments will help to clarify whether the effect of simvastatin can be counteracted by other glucose-lowering agents with mechanisms of action different from sulfonylureas, such as GLP-1 receptor agonists or even SGLT2 inhibitors.

ACKNOWLEDGMENTS

This work was supported by a grant from the University of Padova to AA and FM (Progetto Strategico DYCENDI) and by the EASD Foundation to NE. All named authors meet the International Committee of Medical Journal Editors (ICJME) criteria for authorship for this manuscript, take responsibility for the integrity of the work as a whole, and have given final approval to the version to be published. The article processing charges were funded by the DYCENDI project.

Disclosures. Valentina Scattolini, Camilla Luni, Alessandro Zambon, Silvia Galvanin, Onelia Gagliano, Catalin Dacian Ciubotaru, Angelo Avogaro, Fabio Mammano, Nicola Elvassore, and Gian Paolo Fadini have nothing to disclose.

Compliance with Ethics Guidelines. The institutional review board of the Venetian Institute of Molecular Medicine approved the

study. The protocol followed the Guide for the care and use of laboratory animals, Eighth edition (2011) (<http://grants.nih.gov/grants/olaw/guide-for-the-care-and-use-of-laboratory-animals.pdf>). All institutional and national guidelines for the care and use of laboratory animals were followed.

Data Availability. The datasets used during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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