Efficient transduction of pancreas tissue slices with genetically encoded calcium integrators

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Abstract

This study combines live pancreas tissue slices with adenoviral transduction of the Calcium Modulated Photoactivatable Ratiometric Integrator 2 (CaMPARI2) biosensor for high-throughput analysis of islet calcium responses. Pancreas slices preserve islets within their native microenvironment, adding tissue context to the study of islet function and pathology. A key challenge of the pancreas slice model has been efficient transgene delivery while maintaining viability and function. Here, we demonstrate a robust adenoviral gene delivery approach using targeted and universal promoters. By transducing slices with CaMPARI2 and applying 405 nm photoconverting light, we permanently marked glucose-induced calcium activity across entire islet populations while preserving the in situ tissue context. Applied to nPOD donor tissues, including from individuals with type 1 diabetes, type 2 diabetes, and non-diabetic controls, this approach demonstrated glucose responsive CaMPARI2 labeling that correlated with insulin secretion. Integrating CaMPARI2 with pancreas slices enables multiplexed analyses, linking a functional readout with spatial context through immunostaining or gene expression to advance understanding of human islet behavior.

Introduction

Pancreas tissue slices are an approach to study the physiology of islets and other pancreatic structures in the context of the native tissue microenvironment[1-8]. The slice technique maintains the architecture of the living tissue, including nerves, blood vessels, extracellular matrix, and immune cells. Slices avoid subjecting the pancreatic tissue to the harsh enzymatic and mechanical stress of whole islet isolation. Furthermore, slices allow for the investigation of crosstalk between different pancreatic tissue compartments[9]. Slices make it possible to monitor cell behavior in pathophysiological states that result in compromised islet morphology and where isolation of whole islets is difficult, such as type 1 diabetes (T1D) and type 2 diabetes (T2D)[10].

We and others have successfully used human and mouse pancreas slices to study pancreas physiology in non-diabetic donors and donors with T1D and T2D[10-16]. Functional studies in slices have mostly focused on Ca^{2+} imaging using chemical calcium indicators, with some studies

incorporating electrophysiology as well as the measurement of endocrine hormone secretion and exocrine enzyme release[3,8,10,17-19]. Two studies have used genetically encoded Ca^{2+} indicators (GECI) in slices generated from mice that express these transgenes[11,20].

Robust protocols have been developed for viral transduction in isolated human pancreatic islets and reaggregated pseudo islets, facilitating the use of genetically encoded tools and biosensors[21-26]. Among many potential useful applications are GECI[27,28], optogenetic and chemogenetic actuators[25,29-31], gain or loss of gene function[32], and ex vivo gene editing[33,34]. However, implementing viral gene delivery in human pancreas slices remains challenging. A few examples of adenovirus transduction in slices have been reported. The Domínguez-Bendala group used adenovirus for lineage tracing[35,36] and the Gaisano group demonstrated expression of LC3B-GFP[6]. However, these studies all showed low transduction efficiencies. Thus, consistent, robust viral transduction in pancreas slices has not yet been demonstrated.

Here, we greatly improved adenoviral transduction in pancreatic tissue slices by introducing media agitation and physiological temperature culture, and removing the trypsin inhibitor. This approach substantially increased the number of transduced cells and facilitated the expression of genetically encoded biosensors while maintaining high tissue viability. Functional assays confirmed that transduced tissue slices preserved their responsiveness to stimuli, including real-time analysis of calcium signaling, oxidative stress, and insulin secretion.

We demonstrated the effectiveness of this transduction approach by implementing Calcium Modulated Photoactivatable Ratiometric Integrator 2 (CaMPARI2) to permanently encode Ca^{2+} responses in human pancreas slices. Current Ca^{2+} imaging approaches provide excellent temporal dynamics with real-time imaging of cell behavior including Ca^{2+} responses. However, such approaches are inherently low throughput, where the Ca^{2+} videos are captured one by one. It is typically only possible to obtain functional dynamics for a handful of islets per donor, which greatly under-samples the number of available islets in a slice preparation. Correlating live imaging with later immunostaining in the same islets is technically challenging. Here, we show that a photo-convertible Ca^{2+} integrator can be used to permanently encode Ca^{2+} responses in all islets simultaneously from slice preparations from donors without diabetes and with T1D or T2D. The functional data from these tissues are preserved for correlation with fixed immunostaining. This method provides a valuable tool for investigating questions of functional heterogeneity in the context of the in situ microenvironment for studying disease mechanisms and therapeutic interventions.

Methods

Adenoviral generation

Plasmids, pAV-INS-GRX1-roGFP2 (VB240422-1276nau), and pAV-CMV-CaMPARI2 (VB240306-1476psj) were obtained from VectorBuilder for in-house adenovirus production. The plasmids were received as bacterial glycerol stocks and purified using the ZymoPURE II Plasmid Maxiprep Kit (Zymo Research, D4203-B). Plasmids were first linearized with the PacI restriction enzyme (New England Biolabs, R0547L) and precipitated using 100% isopropanol and 70% ethanol. The linearized DNA was transfected into HEK293T cells in a 6-well plate using Lipofectamine 2000 (Thermo Fisher Scientific, 11668027). Seven days post-transfection, HEK293T cells were examined for fluorescence and viral raft formation. By day 10, cells and media were collected, subjected to three freeze-thaw cycles using a dry ice/ethanol bath to lyse cells, and centrifuged to remove debris. 500 µl of viral supernatant was transferred to a fresh 10 cm plate of HEK293T cells for viral amplification. After two days, infected HEK293T cells

underwent three freeze-thaw cycles for lysis. The adenovirus was subsequently harvested and purified using the Adenovirus Purification Kit (Cell Biolabs, VPK-5112). Adenovirus was then quantified using a QuickTiter Adenovirus Quantification kit (Cell Biolabs, VPK-106). Ad-INS-jRGECO1a and Ad-INS-GRX1-roGFP2 were provided by Dr. Linnemann at Indiana University. Ad-GCaMP6m (Vector Biolabs, 20190319T#9) was purchased from Vector Biolabs in ready-to-use high titer format.

Slice culture and adenoviral transduction

Human pancreas tissue slices were generated by the Network for Pancreatic Organ Donors with Diabetes (nPOD) program at the University of Florida by embedding in agarose and cutting on a vibratome, according to published protocols[10,16,19]. Pancreatic tissue slices used for this study were 120 µm thick, and about 0.5 cm x 0.5 cm in length and width. For insulin secretion studies, pancreatic tissue (1 g) was obtained from Prodo Laboratories and shipped via courier to City of Hope, where slices were generated. After the slices were delivered by nPOD, they were were allowed to recover for 24 hours in an incubator set to 24°C and 5% CO_2 in a 10 cm petri dish with 10 ml of slice culture media consisting of Dulbecco's Low Glucose Modified Eagles Medium +L-Glutamine +Pyruvate (Fisher Scientific SH30021.01), 10% fetal bovine serum (Sigma F4135), 25 kIU/mL aprotinin from bovine lung (Sigma A6106), and 1% antibiotic-antimycotic solution (Corning 30-004-CI). After letting the tissue slices recover for 24 hours, slices were transferred to 35 mm dishes at 2 slices per dish with 2 ml of slice culture media without protease inhibitor. For adenoviral transduction, 5.2×10^8 PFU of ADV (2.6×10^8 PFU per tissue slice) was added to each dish and slices were then cultured statically or on an orbital shaker (Vevor, 0-210) set to 40 rotations per minute inside a 37°C incubator. After 24 hours, viral-containing media was replaced with fresh human slice culture media without protease inhibitor and slices were cultured for an additional 24 hours on the orbital shaker at 37°C before imaging. For secretion studies under different culture conditions, Day 0 control was obtained on the day of slice generation after a 2h resting period on an orbital shaker at room temperature." Slices from Prodo Laboratories were cultured under the same conditions as nPOD slices.

Perifusion

To evaluate glucose-stimulated insulin secretion, three viable pancreatic tissue slices were placed in a closed slice perifusion chamber (Biorep Technologies, PERI-PSC-PP and PERI-PSC-EXT) and connected to a perifusion system with automated tray handling (Biorep Technologies PERI5). The day 0 secretion control was obtained after 2 h resting on an orbital shaker at room temperature, other conditions were cultured for 24 or 48 hours in slice culture media with or without aprotinin and with or without the orbital shaker. Perifusion was performed with HEPES buffer (125 mM NaCl, 5.9 mM KCl, 2.56 mM CaCl2, 1 mM, MgCl2, 25mM HEPES, 0.1% BSA, pH 7.4) containing different glucose concentrations and 2 mM amino acids (L-alanin, L-arginine and L-glutamine (Sigma)). Slices were perfused at a flow rate of 100 µl/min. To remove accumulated hormones and enzymes, pancreatic slices were perfused with HEPES buffer containing 5.5 mM glucose for 90 minutes prior to perfusate collection. Slices were then sequentially exposed to different conditions: 5.5 mM glucose (5G) for 10 minutes and 16.7 mM glucose (16.7G) for 30 minutes. Perfusates were collected at a 60 sec interval in 96 well plates and stored at –20°C until insulin content was measured using a commercially available ELISA kit (Mercodia, human insulin ELISA, 10-1113-01).

Calcium imaging

Live human pancreas tissue slices were transduced using the methods above with the Ca^{2+} biosensor Ad-CMV-GCaMP6m. Live imaging studies were conducted using a Leica SP8 confocal microscope equipped with a live cell incubation system allowing for long-term stable recordings with continual perfusion via an array of programmable syringe pumps (1000-US SyringeONE, New Era Pump Systems, Inc). Slices were initially perfused in Krebs-Ringer Bicarbonate Buffer with HEPES (KRBH, 115 mM NaCl, 4.7 mM KCl, 2.5 mM $CaCl_2$, 1.2 mM KH_2PO_4, 1.2 mM $MgSO_4$, 25 mM $NaHCO_3$, 25 mM HEPES, 0.2% BSA) with 3 mM glucose (3G) then stimulated with KRBH with 16.7mM glucose (16.7G) and finally 3G with carbachol at 10 μM.

Photoconversion of CaMPARI2

Human pancreatic tissue slices were received from nPOD and allowed to recover overnight at 24°C. Tissue slices were transduced with Ad-CMV-CaMPARI2 using the protocol listed above. After the 48-hour transduction period, slices were placed into a 24-well plate with 750 μl of KRBH with 3 mM glucose at 37°C for 10 minutes to allow the slices to equilibrate. An additional 750 μl of 3 mM or 29 mM glucose was then added to the respective groups for 10 minutes before exposing the slices to 405 nm photoconverting (PC) light. Using a 405 nm light-emitting diode (LED) array (Amuza, LEDA-V), the PC light stimulus was delivered using 5 second pulses for 5 mins, leading to 2.5 minutes of total light delivery. The LED array evenly illuminated the entire 24-well plate with an intensity of 400 mW/cm^2. An internal timer using an LED controller (Amuza, LAD-1) was used for precise timing of PC light exposure. For negative control groups, slices were incubated with 3 mM or 16 mM glucose but not exposed to PC light. Control groups were placed into a separate 37°C incubator to avoid any PC light cross-contamination. After PC light exposure, glucose solutions were aspirated, and slices were washed with PBS followed by overnight fixation with 4% paraformaldehyde (PFA) in PBS at 4°C.

Immunohistochemistry

After an overnight fixation in 4% PFA, the slices were washed 3× with PBS and permeabilized with 0.1% Triton X-100 in PBS (TxPBS) for 30 minutes at room temperature (RT). Following permeabilization, the slices were blocked with 10% goat serum in TxPBS for 1 hour at RT to prevent non-specific antibody binding. Primary antibodies diluted in TxPBS with 1% goat serum were applied to the slices and incubated overnight at 4°C. The following day, the slices were washed three times for 5 minutes each with PBS at RT to remove unbound primary antibodies. Secondary antibodies and Hoechst 34580 were then applied and incubated for 1 hour at RT, followed by another three washes with PBS at RT. To prepare for mounting, a circle was drawn on a coverglass using a PAP pen liquid blocker (Newcomer Supply, NC9204359), and 200 μl of PBS was added to facilitate tissue transfer from the 24 well plate. The PBS around the tissue was then aspirated, and a drop of ProLong Gold antifade reagent (Invitrogen, P36930) was applied to the top of the tissue slice. A second coverslip was placed on top, utilizing a double coverslip technique with an imaging spacer (iSpacer, SunJin Lab IS211) to enable imaging from both sides of the tissue slice. The mounted slices were kept at RT overnight to allow the ProLong Gold to set, then stored at 4°C until imaging.

Tile scan analysis

Confocal imaging was performed using a Leica TCS SP8 confocal microscope equipped with a 20×/0.75 NA objective (Leica Microsystems). Tile scans of the entire pancreatic tissue slice were acquired to capture representative images of transduction efficiency throughout the whole tissue. A tile scan with a 10% overlap between adjacent tiles was used to ensure seamless stitching of the individual images. The scan area was defined manually, and the optimal number of tiles was

automatically calculated by the Leica LAS X software. Pinhole size was set to 1 Airy unit for each channel, and the images were acquired at a resolution of 524×524 pixels per tile with a z-step size of 10 µm. For each field of view, a z-stack was acquired. The individual tiles were stitched into a single composite image using Leica LAS X software's automated stitching algorithm. After stitching, images were processed using Fiji (ImageJ) software. Maximum intensity projections of the z-stacks were generated for further quantitative analysis.

Statistics

All data are presented as mean ± SEM. One-way analysis of variance (ANOVA) followed by Tukey's post hoc multiple comparisons test was applied to datasets presented in Figures 4, 5, and 6 to assess group differences. Outlier detection was conducted using the robust regression and outlier removal (ROUT) method with a false discovery rate (Q) set at 1%, identifying two outliers in the 16.7 mM glucose per islet condition for the T2D nPOD case 6619. Area under the curve (AUC) comparisons in Figures 1 and 2 were evaluated using an unpaired two-tailed t-test. Kolmogorov-Smirnov test was used to compare the cumulative distributions in the histograms of Figures 1 and 2.

Reagents

The following primary and secondary antibodies were used for immunofluorescence staining:

Insulin Polyclonal Guinea Pig Antibody (Dako / Agilent no. A0564), CaMPARI Anti-CaMPARI Red 4F6 Mouse Antibody (Absolute Antibody no. AB01649), Goat anti-Mouse IgG (H+L) Highly Cross-Adsorbed Secondary Antibody Alexa Fluor™ 568 (Thermo Fisher no. A11031), Goat anti-Guinea Pig IgG (H+L) Highly Cross-Adsorbed Secondary Antibody Alexa Fluor™ 647 (Thermo Fisher no. A21450).

The following chemicals and reagents were used:

Carbachol (Thermo Fisher, AAL0667403), Dithiothreitol (Sigma, D9779), Hoechst 34580 (Invitrogen, H21486), Isopropanol (Fisher, A416-4), 200 Proof Ethanol (Decon, 2701), TE Buffer (Fisher, BP2473100), FBS (Gibco, 10082147), Sodium bicarbonate (Sigma, S5761), Sodium chloride (Sigma, S5886), PBS (Thermo Fisher Scientific, 20012050), HEPES (Sigma, H4034), HBSS (Gibco, 14025092), Penicillin Streptomycin (Gibco, 15140122), Potassium phosphate monobasic (Sigma, P5655), Magnesium sulfate heptahydrate (Sigma, M2773), BSA (Fisher Scientific, 199898), Calcium chloride (Sigma, C5670), Potassium chloride (Sigma, P5405), D-(+)-Glucose (Sigma, G7021), Paraformaldehyde (Sigma, P6148), Tolbutamide (Sigma, T0891), Diazoxide (Sigma, D9035), Forskolin (Sigma, F6886).

Study approval

All experimental protocols using mice were approved by the University of Florida Animal Care and Use Committee (201808642, 202109991, 202400000147). All procedures using human slices were performed according to the established standard operating procedures of the nPOD/OPPC and approved by the University of Florida Institutional Review Board (IRB201600029) and the United Network for Organ Sharing (UNOS) according to federal guidelines, with informed consent obtained from each donor's legal representative. Pancreas tissue from human donors with or without T1D was procured from the nPOD program at the University of Florida (RRID: SCR_014641, https://www.jdrfnpod.org). For each donor, a medical chart review was performed, and C-peptide was measured, with T1D diagnosed according to the

guidelines established by the American Diabetes Association (ADA). Demographic data, hospitalization duration, and organ transport time were obtained from hospital records. Donor pancreata were recovered, placed in transport media on ice, and shipped via organ courier to the University of Florida. The tissue was processed by a licensed pathology assistant. Detailed donor information is listed in Table 1. nPOD tissues specifically used for this project were approved as nonhuman by the University of Florida IRB (NH00041892, NH00042022).

Results

Transduction on orbital shaker improves transduction efficiency

Robust expression of genetically encoded tools and biosensors for studying cellular processes in has been challenging to achieve in pancreas tissue slices. Here, we modified pancreas slice culture procedures to develop an approach for efficient viral gene delivery. Pancreas tissue slices secrete the digestive enzymes amylase, lipase, and trypsinogen, which can compromise slice viability[19]. To mitigate pancreatic enzyme activity, human pancreas slices are typically cultured with aprotinin at 28-30°C[10,16,19]. The lower culture temperature, combined with the use of trypsin inhibitor, prevents self-digestion and slows metabolism, thereby extending slice viability. These culture conditions also seem to hinder efficient viral transduction[37]. Additionally, in a static culture environment, transduction is limited by reduced viral particle collisions, as large particles like virions diffuse more slowly than small solutes[38]. We hypothesized that introducing media agitation would enhance viral mass transport to the tissue slices. Unlike monolayer cultures, where cells are evenly distributed, cells in tissue slices are highly concentrated in a small area. We hypothesized that agitation would help disperse the secreted pancreatic enzymes more efficiently than diffusion alone. This elevated efficiency should then allow for slice culture at 37°C without protease inhibitor during the brief viral transduction period.

To test this hypothesis, we initially focused on delivering genetically encoded calcium indicators (GECI) to slices, as these biosensors allow us to assess islet cell function post-transduction by Ca^{2+} imaging. To improve pancreas tissue slice transduction, we removed protease inhibitor from the media, increased the culture temperature to 37°C, introduced media agitation by orbital shaker, and used a very high viral particle concentration of 5.2×10^8 PFU/ml. A virus concentration of 1×10^6 PFU/ml is typical for transducing cell monolayers. A standard viral titer for transducing cultured cells appears to be insufficient for good transduction of organotypic slices. Previous studies using adenovirus to transduce pancreas slices reported an MOI of 10 to 50 [35,36]. We used an MOI of 86 in our protocol, assuming a typical pancreas slice is 5,000 µm x 5,000 µm x 120 µm in dimension and contains around 3×10^6 cells, with two slices per dish in 2 ml of transduction media.

We initially compared the transduction of pancreas slices with and without media agitation, keeping MOI, lack of aprotinin, and temperature the same in both groups. At 24 hours, media was exchanged for fresh virus-free media, and we waited an additional 24 hours for transgene expression to occur. Transduction efficiency was quantified by taking confocal tile scans of the entire tissue slice and analyzing the mean fluorescence intensity (MFI) of the maximum intensity projection of each tissue slice (n=5 tissue slices per group) (Figure 1A). We observed low fluorescence and sporadic transduction in the statically cultured slices but widespread and bright fluorescence in the slices with media agitation. The simple addition of the low-speed orbital shaker during culture increased the mean fluorescence intensity of GCaMP6m by approximately 3x (Figure 1B).

While higher temperature culture, removal of protease inhibitor, and media agitation using an orbital shaker greatly improved transduction efficiency, these conditions could potentially be detrimental to islet function. To investigate any detrimental effects, we conducted a dynamic perifusion assay to evaluate insulin secretion from pancreatic tissue slices cultured under orbital shaker conditions compared to static culture after 24 and 48 hours and with and without the protease inhibitor aprotinin (apr) (Figure 1C). As measured by the area under the curve (AUC) during the stimulation phase, slices cultured on the orbital shaker at 37°C for 24 or 48 hours and with or without aprotinin were not statistically different from slices tested on the day of generation (day 0) or from slices cultured under static conditions with or without protease inhibitor (Figure 1D). These data indicate that addition of orbital culture and removal of protease inhibitor were not detrimental to glucose responsive insulin secretion in slices. As human slices are a limited resources with substantial variability, we pooled the AUC datapoints from the different static and orbital groups to increase the sample size and compared static versus orbital head-to-head. In this comparison, orbital culture resulted in significantly improved insulin secretion over static culture (Figure 1E).

Endocrine and exocrine function retained post adenoviral transduction

Adenoviral transduction in dense tissues often requires high multiplicities of infection (MOIs), which can result in cytotoxicity. While the tissue may express the biosensor, cellular responsiveness may be compromised. In this study, we demonstrate by using the Ca^{2+} biosensor Ad-CMV-GCaMP6m, high transduction efficiency can be achieved while maintaining both endocrine and exocrine cell responsiveness 72 hours post-adenoviral transduction when cultured on an orbital shaker. Human pancreas slices from a donor without diabetes transduced with the GECI Ad-CMV-GCaMP6m under either orbital or static conditions were assessed for dynamic Ca^{2+} responses to 16.7 mM glucose and the cholinergic agonist carbachol. Representative snapshots of peak calcium influx are shown in Figure 2A, with heatmaps and dynamic perfusion traces of calcium responses presented in Figures 2B and 2C. Of note, fewer transduced cells were visible in the static culture group due to lower transduction efficiency and expression levels. Slices transduced under orbital shaker conditions had a significantly greater area of cells expressing the probe and brighter probe fluorescence within transduced cells (Figure 2E). Tissue slices transduced under orbital shaking conditions exhibited greater calcium influx in response to 16.7 mM glucose in endocrine cells and to 10 μM carbachol in exocrine cells, compared to slices transduced in static culture (n=3) (Figure 2D) (Supplementary Video 1).

Targeted expression in specific cell types

An advantage of genetically expressed biosensors over chemical sensors is the ability to target specific cell types using cell type specific promoters. In pancreatic tissue slices, where multiple cell types are present, insulin promoters ensure that data is collected only from beta cells. Here we demonstrate that we can target pancreatic beta cells using a reactive oxygen species (ROS) sensor driven by a hybrid insulin promoter (RIP / rabbit BGI)[39], Ad-INS-GRX1-roGFP2 with minimal leaky expression (Figure 3A). We also show the ability to co-express multiple targeted sensors for multiplexed assays, using both a Ca^{2+} biosensor, Ad-INS-jRGECO1a, and Ad-INS-GRX1-roGFP2 (Figure 3B). GRX1-roGFP2 is a fusion of human glutaredoxin-1 (GRX1) to redox-sensitive GFP (roGFP2) that facilitates live imaging of the glutathione redox potential (E(GSH)) with high temporal resolution[39,40]. jRGECO is a red fluorescent protein-based GECI[41]. Following transduction, the function of the ROS probe and tissue responsiveness was confirmed by perfusing pancreatic tissue slices with hydrogen peroxide (H_2O_2). Upon 20 μM H_2O_2 exposure, an

increase in the oxidation ratio of Ad-INS-GRX1-roGFP2 was observed, indicating successful detection of ROS. To further confirm the functionality and reversibility of the sensor, the tissue slices were subsequently treated with 1 mM dithiothreitol (DTT), a reducing agent (Figure 3C). This treatment successfully restored the baseline oxidation ratio, demonstrating that the ROS sensor retained its sensitivity and reversibility post-transduction. These data demonstrate the ability to specifically transduce beta cells in pancreas slices and multiplex multiple probes in a single slice.

High throughput screening of beta cell glucose responsiveness in tissue slices

Calcium Modulated Photoactivatable Ratiometric Integrator 2 (CaMPARI2) is a photoconvertible protein that enables permanent marking of Ca^{2+} activity in large populations of cells over defined time windows. CaMPARI2 shifts from green to red emission upon simultaneous irradiation with violet light (405 nm) in the presence of elevated intracellular calcium [42,43]. CaMPARI2 enables the permanent marking of cells that have experienced an increase in intracellular Ca^{2+} during a defined time window of photoconverting (PC) light exposure. Simultaneous PC light and elevated Ca^{2+} are required for photoconversion of the probe.

Real-time calcium imaging in human pancreatic tissues has traditionally relied on chemical sensors, such as Fluo-4, confined to the microscope's live field of view. Here, we demonstrate the use of adenoviral transduction of pancreatic tissue slices with a genetically encoded Ca^{2+} integrator, Ad-CMV-CaMPARI2, allowing for high throughput calcium activity assessment by encoding total calcium activity across the entire tissue during a defined stimulation window. This approach facilitated the simultaneous functional assessment of all the islets across up to 24 slices from a single nPOD slice case in one experiment (~60 islets). No microscope was involved during the stimulation and photoconversion protocol. Calcium activity was only quantified after-the-fact in fixed and immunostained slices by measuring the photoconversion within insulin positive cells.

First, we assessed a slice donor with no diabetes to confirm the function of CaMPARI2 in our system. CaMPARI2 requires the simultaneous presence of elevated Ca^{2+} and PC light to convert from green to red states. Live slices transduced with Ad-CMV-CaMPARI2 using the orbital shaker method were placed in a 24 well plate (1 slice per well) and statically exposed to 3 mM or 16 mM glucose with or without PC light from a 405 nm light-emitting diode array positioned directly under the cell culture plate. Following PC light exposure, slices were fixed and immunostained for insulin and with antibody that specifically labels only the photoconverted red form of CaMPARI2 and not the unconverted green form of CaMPARI2[43]. We used this monoclonal antibody to strongly label any cells that had calcium activity during the stimulation protocol. Fixed, stained, and mounted slices were imaged by confocal at a time of the investigator's choosing, which neutralized much of the uncertainty and randomness surrounding when nPOD slice donor cases arrived and allowed for careful and meticulous observations of all islets in the sample.

By confocal imaging of the fixed and immunostained slices, no photoconversion was observed in the CaMPARI2 expressing slices that were not exposed to PC light. There was no photoconversion for either 3 mM or 16 mM glucose without PC light, demonstrating no spontaneous photoconversion of CaMPARI2 (Figure 4A). Low amounts of photoconversion were seen in 3 mM glucose with PC light, reflecting basal islet calcium activity. In contrast, strong photoconversion was observed with the combination of 16 mM glucose and PC light, confirming that beta cells responded to glucose stimulation by mobilizing cytosolic calcium and this response was encoded and preserved in the fixed slices (Figure 4A). We quantified the CaMPARI2 photoconversion in beta cells by hand-drawing regions of interest (ROIs) over cells double-positive for CaMPARI2 and insulin and measuring the ratio of mean fluorescence intensity of

CaMPARI2 red to CaMPARI2 green on a per beta cell and per islet basis (Figure 4B,C). Interestingly, we observed significant heterogeneity in beta cell responses to glucose at both the single beta cells and single islet level. Despite an overwhelmingly clear response to glucose, two of the nine islets from this non-diabetic donor did not respond to 16 mM glucose, suggesting CaMPARI2 can be used to investigate islet functional heterogeneity.

Having demonstrated that the probe is functional for encoding calcium activity in beta cells within human pancreatic tissue slices, we aimed to investigate whether we could use this probe to assess beta cell function in various diseased states. We compared beta cell function in a second donor without diabetes, a donor with T2D, and a donor with T1D. Donor characteristics are reported in Table 1. The donor without diabetes, (nPOD case 6622) showed strong glucose responsive Ca^{2+} mobilization when assessed by photoconversion ratio of the CaMPARI2 probe. The beta cells displayed strong calcium activity upon high glucose and KCl stimulation, resulting in significant CaMPARI2 photoconversion (Figure 5A, 5B), consistent with dynamic insulin secretion data from this case (Figure 5C).

Repeating the same experimental design for the calcium assay in a donor with T2D (nPOD case 6619), we observed a dysregulation in beta cell glucose sensing, particularly elevated CAMPARI2 conversion in low glucose conditions. (Figure 6A, 6B). This finding is consistent with insulin perifusion data, which also showed elevated insulin secretion at low glucose levels (Figure 6C). Additionally, the fold change in photoconversion between low and high glucose was markedly lower compared to the non-diabetic case, further highlighting the impaired glucose responsiveness in the beta cells of this particular T2D case, a phenotype that has been observed in other slices from donors with T1D[44].

The CaMPARI2 assay was also performed in slices from a donor with T1D (nPOD case 6618), which had a disease duration of 2 years. Despite the significantly reduced beta cell mass in this case, we successfully captured the calcium activity from two remaining insulin-positive islets, one each under low and high glucose conditions. The residual beta cells were non-functional, showing no response to high glucose stimulation, resulting in low photoconversion of CaMPARI2 (Figure 7A, 7B). These findings are consistent with perifusion data for this case, which demonstrated a total absence of insulin release in response to both high glucose and KCl stimulations (Figure 7C). This data is also consistent with beta cell glucose blindness observed in live Ca^{2+} imaging in pancreas slices from donors with T1D[10,16,19].

Overall, our findings demonstrate the utility of CaMPARI2 as a biosensor for permanently encoding calcium activity in human pancreatic tissue slices during a defined stimulation window. This tool can be implemented across various disease states, including T1D and T2D, as well as across different stages of disease progression, offering a powerful, higher throughput means to assess calcium dynamics and glucose responsiveness.

Discussion

Adenoviral transduction in pancreatic tissue slices presents a unique challenge as they continuously secrete digestive enzymes, which can degrade viral particles and impact the viability of the tissue. Additionally, as opposed to transduction of cell monolayers, where cells are evenly spread out across the culture dish, cells within a tissue slice are densely confined to a small area, reducing the probability of viral particle collisions. A study of adenoviral transduction of human brain slices used microinjection of high titer virus directly into the slice tissue[45]. In static culture conditions commonly used for transduction, viral particles diffuse more slowly than smaller molecules, reducing their interactions with the tissue[38]. Our study demonstrates that gentle media

agitation during adenoviral transduction significantly improves transduction efficiency compared to static conditions, without compromising tissue functionality. The increased transduction efficiency, evidenced by a 300% rise in MFI, suggests that media agitation facilitates the movement of viral particles, overcoming diffusion limitations and improving the chances of viral particles reaching the small surface area of pancreatic tissue slices. We anticipate that our approach will be conducive to transducing organotypic slices from different tissues and with other virus types including lentivirus and adeno-associated virus.

The potential limitation of orbital shaking is the introduction of mechanical stress, which could adversely affect beta cell function. To address this concern, we evaluated the functional integrity of transduced tissue slices through dynamic perifusion assays. Our results demonstrated that pancreas slices cultured under mild orbital shaking conditions not only retained but enhanced their insulin secretion capacity in response to glucose. This suggests that gentle orbital shaking does not negatively impact beta cell function; instead, it may contribute to maintaining tissue viability and responsiveness.

In addition to enhanced transduction efficiency, we showed that adenoviral transduction does not compromise the functional responsiveness of either endocrine or exocrine cells using real time calcium imaging. Calcium imaging using the genetically encoded calcium indicator Ad-CMV-GCaMP6m revealed strong calcium responses to glucose in islets and to carbachol in exocrine cells, further highlighting the preserved functionality of the transduced tissue. Tissue slices transduced under orbital shaking conditions exhibited a greater number of transduced cells and stronger calcium responses compared to static culture. A variety of probes can be used in slices, targeted to specific cell types, and even multiplexed, as shown by our demonstration of Grx1-roGFP2 and jRGECO1a.

The use of the genetically encoded calcium integrator CaMPARI2 allowed us to conduct high throughput calcium assays, facilitating the functional assessment of all the islets in our allocation of slices from an nPOD donor case. The transient nature of traditional calcium indicators, such as Ad-CMV-GCaMP6m used in Figure 3, provide a limited field of view and require continuous monitoring islets to capture peak calcium activity upon glucose stimulation. Selecting which islets to record can be biased, resulting in data that is not reflective of the entire islet population in the tissue. All calcium recordings within a case are typically performed on the same day to minimize variability, but due to the time required to set up and run each recording, only 3 to 4 islets per case are usually captured. This constraint highlights the advantages of genetic tools like CaMPARI2 in pancreatic tissue slices. By enabling the simultaneous analysis of entire islet populations, CaMPARI2 reduces islet selection bias and ensures that data is reflective of the whole population. This is especially critical in rare cases, such as donors with T1D, where samples may only be available sporadically. Having a tool like CaMPARI2 ensures we maximize the value of the sample, increasing the likelihood of capturing representative data.

Efficient protocols for adenoviral transduction in slices enabled us to examine calcium dynamics across different disease states, including T1D and T2D cases. Our findings revealed dysregulated glucose sensing in T2D case 6619, characterized by elevated calcium influx under low glucose conditions, consistent with abnormal insulin secretion observed in perifusion assays. In T1D case 6618, we observed non-functional beta cells that failed to respond to high glucose stimulation, reflecting the loss of glucose-responsive insulin secretion in this case. The ND case 6622 showed preserved beta cell function, as evidenced by normal calcium activity and insulin secretion responses to glucose stimulation. While these findings are promising, they are based on a single case each. In this paper, we demonstrate the feasibility of using this tool to monitor calcium dynamics in various states, but further studies with more cases are needed to establish

meaningful patterns or differences across disease states. Nonetheless, our work shows that CaMPARI2 is a powerful biosensor for detecting changes in calcium activity in beta cells within human pancreatic tissue slices across various disease stages.

Our findings also revealed significant heterogeneity in beta cell calcium responses within individual islets. This variation may reflect functional subpopulations of beta cells, such as first-responder or leader cells, which have been proposed to initiate coordinated islet activity. Studies have identified leader cells as highly responsive beta cells that drive islet-wide calcium waves[46-48]. Future studies using CaMPARI2 in conjunction with functional and transcriptomic profiling could help delineate the molecular identity of these leader cells and their role in glucose regulation.

In future work, we plan to use CaMPARI2 to correlate various functional and structural parameters within the islets of tissue slices. For instance, using this tool we can explore the relationship between calcium responsiveness and islet architecture, such as the size, morphology, or location of the islet. Additionally, integrating CaMPARI2 with spatial transcriptomics could allow us to link functional calcium responses with gene expression patterns in specific islet regions, providing a molecular context to observed functional heterogeneity. Furthermore, CaMPARI2 can be combined with immune cell staining techniques to quantify islet function during insulitis, offering insights into how immune cell presence might impact beta cell function and survival. These capabilities open opportunities for understanding complex dynamics within pancreatic tissue slices, particularly in diseases like T1D, where both functional and immune-mediated factors are critical.

Conclusions

Our study establishes an optimized adenoviral transduction protocol for human pancreatic tissue slices, significantly enhancing transgene delivery while preserving tissue viability and function. By incorporating media agitation, higher temperature culture, and removal of trypsin inhibitors during transduction, we significantly improved transduction efficiency without compromising endocrine or exocrine cell responsiveness. These optimizations address a critical technical limitation in pancreas slice research, facilitating the application of genetically encoded biosensors and functional genetic manipulations in human pancreatic tissue.

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Author contributions

CSL developed the methodology for culturing and transducing live human pancreatic tissue slices on an orbital shaker, conducted fixed and real-time Ca^{2+} imaging studies, performed fixed tissue staining and imaging, analyzed Ca^{2+} imaging and CaMPARI2 data, generated adenovirus, and drafted the manuscript. AES designed the setup for CaMPARI2 experiments, assisted with CaMPARI2 experiments, contributed to the development of imaging approaches, and generated adenovirus. JKP performed perifusion studies and insulin ELISAs on human tissue slices. AKL provided critical reagents and technical assistance. MB and HH generated live human pancreatic tissue slices and conducted perifusion studies. CLS and CEM provided critical funding, guidance and technical assistance. EAP secured funding, conceived the study, supervised the research, and edited the manuscript. All authors contributed to data interpretation, reviewed and commented on the manuscript, and approved its final submission.

Table 1

Case ID	Donor Type	Age (years)	Sex	BMI (kg/m²)	Figure
6558	Single autoantibody positive (GADA)	21	Female	27.8	1A, 1B
6559	No Diabetes	23	Female	26.3	2
6548	No Diabetes	20	Male	23.8	3
6617	No Diabetes	19	Male	22.8	4
6618	Type 1 Diabetes (2 years duration)	31	Male	23.6	5
6619	Type 2 Diabetes	14	Male	37.3	6

Table 2

Case ID	Donor Type	Age (years)	Sex	BMI (kg/m²)	Figure
HP23241	No Diabetes	59	Male	28.1	1C,1D
HP23270	No Diabetes	34	Male	27.0	1C,1D
HP23285	No Diabetes	58	Female	21.0	1C,1D
HP23305	No Diabetes	47	Male	28.3	1C,1D

Figure Legends

Figure 1: Orbital Shaker vs. Static Transduction and Insulin Secretion. (A) Confocal tilescan images of Ca^{2+} sensor Ad-CMV-GCaMP6m expression in whole human pancreatic tissue slices (nPOD 6558), transduced on an orbital shaker or in static culture with no aprotinin. Slices were transduced using 2.6×10^8 PFU of Ad-CMV-GCaMP6m per tissue slice, incubated for 24 hours at 37°C, and imaged 48 hours post-transduction. **(B)** Transduction efficiency was quantified using Mean Fluorescence Intensity (MFI), (± SEM, n = 5, students unpaired t-test, ** $p < 0.01$). **(C)** Dynamic insulin secretion of human pancreatic tissue slices from three to four nondiabetic donors were cultured on an orbital shaker (blue) or in static culture (red) after 24 and 48 hours, with and without aprotinin (apr). Insulin secretion traces from slices on Day 0 (gray) is repeated on all plots as a control. Slices were initially perfused with 5.5 mM glucose (5G) for 10 minutes, followed by stimulation with 16.7 mM glucose (16.7G) for 30 minutes. Data are normalized to the average baseline secretion at 5.5G. **(D)** Quantification of insulin secretion from perifusion assays in (C), shown as the area under the curve (AUC) for 16.7G stimulation at 0, 24 and 48 hours with or

without aprotinin. AUC data from slices on Day 0 (gray) are provided as a control. **(E)** Same AUC data as in (D), but with aprotinin and no-aprotinin groups combined at 24 and 48 hours. (± SEM, n = 18, students unpaired t-test, * $p < 0.05$). The data are representative of the average of three to four donors, with each donor contributing four slices.

Figure 2: Endocrine and Exocrine Function Retained 72 Hours Post Adenoviral Transduction. Realtime confocal time-lapse perfusion study of pancreatic tissue slices (nPOD 6559) expressing the adenoviral Ca^{2+} biosensor CMV-GCaMP6m, transduced using an orbital shaker (bottom row) or static culture (top row). Endocrine and exocrine tissues were stimulated using perfusion of either 16.7 mM high glucose (16.7G) or 10 μM carbachol, respectively. **(A)** Images were selected from the real-time perfusion study at the peak Ca^{2+} influx of the 16.7 mM stimulation, islets outlined in white. **(B)** Heat maps of realtime Ca^{2+} responses in individual cells in both islet and acinar tissues shown in (A), imaged 48 hours post-transduction. Baseline activity was first acquired by perfusion of low 3 mM glucose (3G) for five minutes, followed by 16.7G, eliciting Ca^{2+} influx within the pancreatic islet. High glucose was washed away by perfusion of 3G, followed by stimulation of acinar tissue using 10 μM carbachol. **(C)** Average real-time Ca^{2+} traces of (n=3) slices following the protocol outlined in (B). The X-axis is segmented to separately show Ca^{2+} influx in the islet (left) and acinar tissues (right). **(D)** Quantification of the Ca^{2+} traces in (C), shown as the area under the curve (AUC) for 16.7G stimulation in the islet or 10 μM carbachol stimulation in the acinar tissue. (± SEM, n = 5, students unpaired t-test, * $p < 0.05$, ** $p < 0.01$) **(E)** Histograms depicting the sum of pixel intensity values at peak Ca^{2+} influx in the islet during high glucose stimulation and in the acinar tissue during carbachol stimulation. (Kolmogorov-Smirnov nonparametric cumulative distributions test, **** $p < .0001$).

Figure 3: Targeted expression in specific cell types. Confocal images of beta cell-targeted expression of adenoviral vectors INS-GRX1-roGFP2 (green) and INS-jRGECO1a (red). **(A)** Representative image of a human pancreas slice alongside higher magnification images of islets within the tissue (nPOD 6548). Slices were transduced using 2.6×10^8 PFU of INS-GRX1-roGFP2 per tissue slice, incubated for 24 hours at 37°C, and imaged 48 hours post-transduction. **(B)** Dual expression of INS-GRX1-roGFP2 (green) and INS-jRGECO1a (red) in an islet within a human pancreas slice. Dual expression was achieved in slices using 1.4×10^8 PFU of INS-GRX1-roGFP2 and 1.4×10^8 PFU of INS-jRGECO1a per tissue slice, incubated for 24 hours at 37°C. Confocal images were taken 48 hours post-transduction. **(C)** Perfusion of human pancreatic tissue slice in a Warner imaging chamber using 3 mM glucose for 3 minutes, followed by a 20 μM H2O2 stimulation. ROS detoxification was achieved by perfusing 1 mM DTT at 12 minutes.

Figure 4: Functional Ca^{2}+ Biosensor, CaMPARI2, in Living Human Pancreas Tissue Slices (nPOD 6617 ND). Cells that respond to glucose stimulation experience an influx of calcium ions. An increase in intracellular calcium, along with exposure to ultraviolet (UV) light for photoconversion (PC), irreversibly converts the fluorescence of CaMPARI2 from green to red. **(A)** Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in high or low glucose for ten minutes, followed by exposure or no exposure to PC light. PC light-treated groups were exposed using ten second pulses for five minutes. Tissue slices were then chemically fixed with paraformaldehyde for staining. **(B, C)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 red:green ratio. Only cells that were positive for insulin were quantified (n = 3 slices per group). Data were analyzed per beta cell or per islet. (± SEM, one-way ANOVA with Tukey's post-hoc test, *** $p < 0.001$, **** $p < 0.0001$, outliers removed with ROUT Q = 1%).

Figure 5: Functional Ca^{2+} Biosensor, CaMPARI2, in Living Human Pancreas Tissue Slices (nPOD 6622 ND Control). (A) Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in 3 mM and 16.7 mM glucose for ten minutes, or 30 mM KCl for one minute. Slices were

then exposed to PC light using ten second pulses for five minutes before being washed with PBS. Tissue slices were then chemically fixed with paraformaldehyde for staining. Beta cells displayed strong calcium activity upon high glucose stimulation, resulting in significant photoconversion. **(B)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 Red:Green Ratio. Only cells that were positive for insulin were quantified. (n = 4) slices per group. (± SEM, one-way ANOVA with Tukey's post-hoc test, *** $p < 0.001$, **** $p < 0.0001$, outliers removed with ROUT Q = 1%). **(C)** Perifusion analysis of pancreatic tissue slices from this non-diabetic donor demonstrated robust insulin secretion in response to high glucose stimulation, confirming the preserved glucose responsiveness of beta cells.

Figure 6: Functional Ca^{2+} Biosensor, CaMPARI2, in T2D nPOD Case 6619. (A) Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in 3 mM and 16.7 mM glucose for ten minutes or 30 mM KCl for one minute. Slices were then exposed to photoconversion (PC) using ten second pulses for five minutes before washing with PBS. Tissue slices were then chemically fixed with paraformaldehyde for staining. **(B)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 Red:Green ratio. We observed a dysregulation in beta cell glucose sensing, particularly elevated calcium influx in low glucose conditions. Only insulin-positive cells were quantified (n = 4 slices per group). Data were analyzed on a per-beta-cell or per-islet basis. (± SEM, one-way ANOVA with Tukey's post-hoc test, *** $p < 0.001$, **** $p < 0.0001$, outliers removed with ROUT Q = 1%). **(C)** The T2D case exhibited an atypical response, with significantly elevated insulin secretion at low glucose levels, aligning with the dysregulated calcium influx observed using CaMPARI2.

Figure 7: Functional Ca^{2+} Biosensor, CaMPARI2, in T1D nPOD case 6618. (A) Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in 3 mM and 16.7 mM glucose for ten minutes. Slices were then exposed to PC light using ten second pulses for five minutes before washing with PBS. Tissue slices were then chemically fixed with paraformaldehyde for staining. **(B)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 red:green ratio. The residual beta cells were non-functional, showing no response to high glucose stimulation, resulting in no photoconversion of CaMPARI2. Only cells that were positive for insulin were quantified (n = 1 slice per group). Data were analyzed on a per-beta-cell basis. **(C)** Perifusion analysis of pancreatic tissue slices from this T1D case revealed a total absence of insulin release in response to both high glucose and KCl stimulations, confirming the non-functionality of the remaining beta cells (red). Perifusion trace from non-diabetic (ND) case 6622 (grey) provided as a control.

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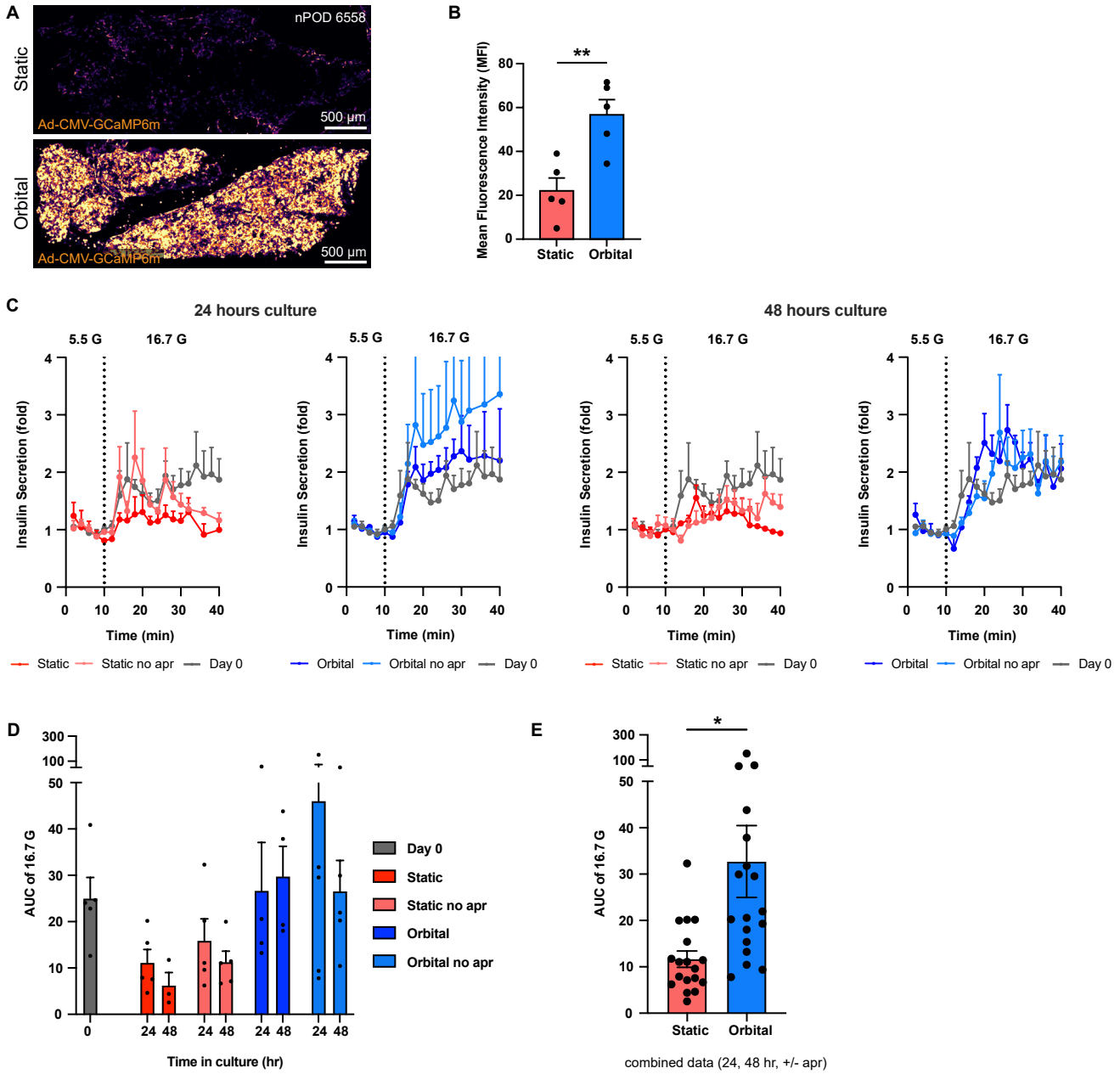

Figure 1: Orbital Shaker vs. Static Transduction and Insulin Secretion. (A) Confocal tilescan images of Ca^{2+} sensor Ad-CMV-GCaMP6m expression in whole human pancreatic tissue slices (nPOD 6558), transduced on an orbital shaker or in static culture with no aprotinin. Slices were transduced using 2.6 × 10^8 PFU of Ad-CMV-GCaMP6m per tissue slice, incubated for 24 hours at 37°C, and imaged 48 hours post-transduction. **(B)** Transduction efficiency was quantified using Mean Fluorescence Intensity (MFI), (± SEM, n = 5, students unpaired t-test, ** $p < 0.01$). **(C)** Dynamic insulin secretion of human pancreatic tissue slices from three to four nondiabetic donors were cultured on an orbital shaker (blue) or in static culture (red) after 24 and 48 hours, with and without aprotinin (apr). Insulin secretion traces from slices on Day 0 (gray) is repeated on all plots as a control. Slices were initially perfused with 5.5 mM glucose (5G) for 10 minutes, followed by stimulation with 16.7 mM glucose (16.7G) for 30 minutes. Data are normalized to the average baseline secretion at 5.5G**. (D)** Quantification of insulin secretion from perifusion assays in (C), shown as the area under the curve (AUC) for 16.7G stimulation at 0, 24 and 48 hours with or without aprotinin. AUC data from slices on Day 0 (gray) are provided as a control. **(E)** Same AUC data as in (D), but with aprotinin and no-aprotinin groups combined at 24 and 48 hours. (± SEM, n = 18, students unpaired t-test, * $p < 0.05$). The data are representative of the average of three to four donors, with each donor contributing four slices.

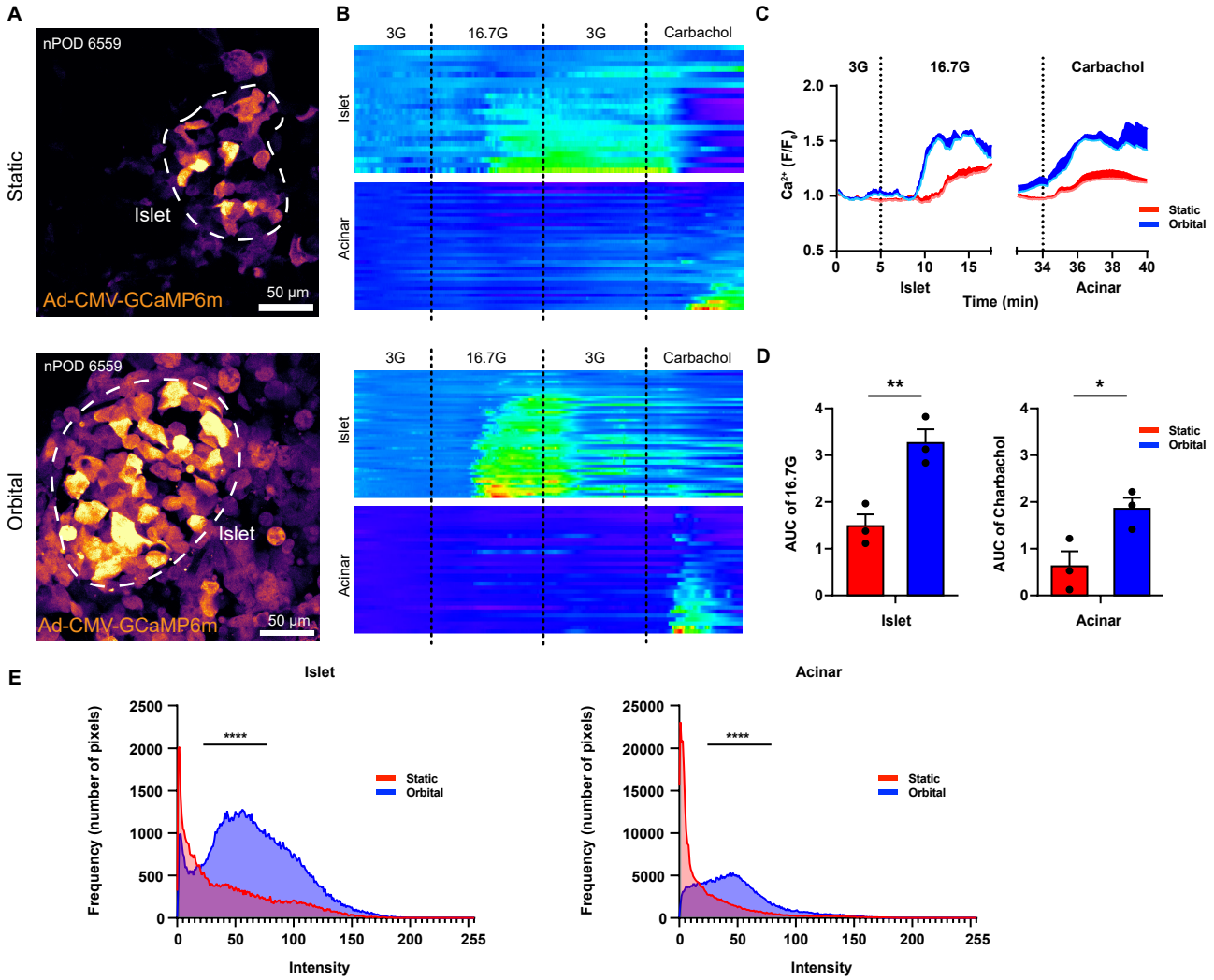

Figure 2: Endocrine and Exocrine Function Retained 72 Hours Post Adenoviral Transduction. Real-time confocal time-lapse perfusion study of pancreatic tissue slices (nPOD 6559) expressing the adenoviral Ca^{2+} biosensor CMV-GCaMP6m, transduced using an orbital shaker (bottom row) or static culture (top row). Endocrine and exocrine tissues were stimulated using perfusion of either 16.7 mM high glucose (16.7G) or 10 µM carbachol, respectively. **(A)** Images were selected from the real-time perfusion study at the peak Ca^{2+} influx of the 16.7 mM stimulation, islets outlined in white. **(B)** Heat maps of realtime Ca^{2+} responses in individual cells in both islet and acinar tissues shown in (A), imaged 48 hours post-transduction. Baseline activity was first acquired by perfusion of low 3 mM glucose (3G) for five minutes, followed by 16.7G, eliciting Ca^{2+} influx within the pancreatic islet. High glucose was washed away by perfusion of 3G, followed by stimulation of acinar tissue using 10 µM carbachol. **(C)** Average real-time Ca^{2+} traces of (n=3) slices following the protocol outlined in (B). The X-axis is segmented to separately show Ca2+ influx in the islet (left) and acinar tissues (right). **(D)** Quantification of the Ca^{2+} traces in (C), shown as the area under the curve (AUC) for 16.7G stimulation in the islet or 10 µM carbachol stimulation in the acinar tissue. (± SEM, n = 5, students unpaired t-test, * $p < 0.05$, ** $p < 0.01$) **(E)** Histograms depicting the sum of pixel intensity values at peak Ca^{2+} influx in the islet during high glucose stimulation and in the acinar tissue during carbachol stimulation. (Kolmogorov-Smirnov nonparametric cumulative distributions test, **** $p < .0001$).

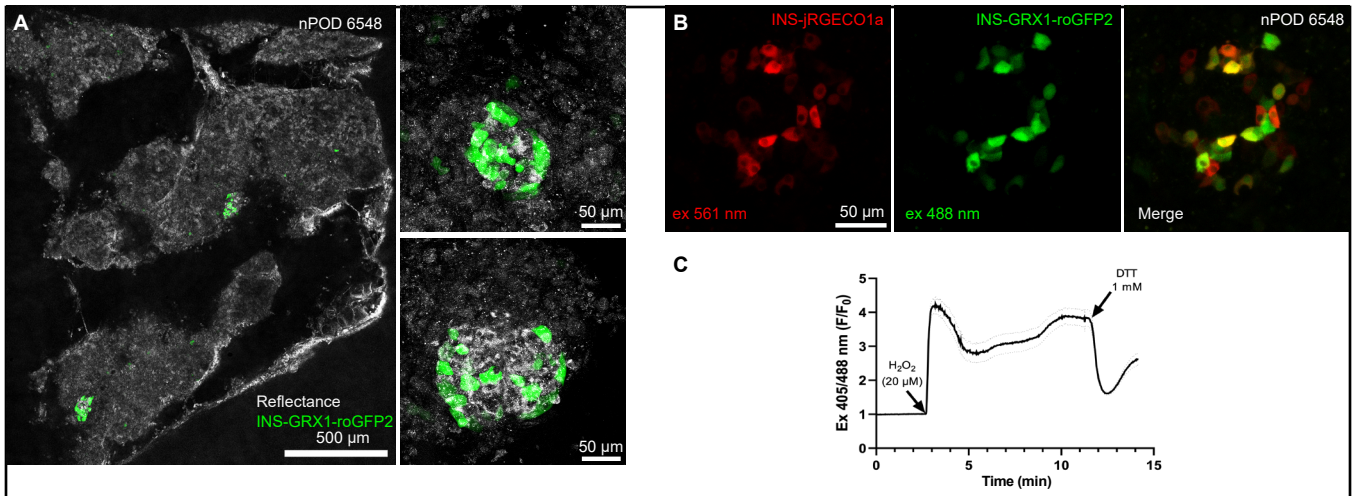

Figure 3: Targeted expression in specific cell types. Confocal images of beta cell-targeted expression of adenoviral vectors INS-GRX1-roGFP2 (green) and INS-jRGECO1a (red).
(A) Representative image of a human pancreas slice alongside higher magnification images of islets within the tissue (nPOD 6548). Slices were transduced using 2.6 × 10^8 PFU of INS-GRX1-roGFP2 per tissue slice, incubated for 24 hours at 37°C, and imaged 48 hours post-transduction. **(B)** Dual expression of INS-GRX1-roGFP2 (green) and INS-jRGECO1a (red) in an islet within a human pancreas slice. Dual expression was achieved in slices using 1.4 × 10^8 PFU of INS-GRX1-roGFP2 and 1.4 × 10^8 PFU of INS-jRGECO1a per tissue slice, incubated for 24 hours at 37°C. Confocal images were taken 48 hours post-transduction. **(C)** Perfusion of human pancreatic tissue slice in a Warner imaging chamber using 3 mM glucose for 3 minutes, followed by a 20 µM H_2O_2 stimulation. ROS detoxification was achieved by perfusing 1 mM DTT at 12 minutes.

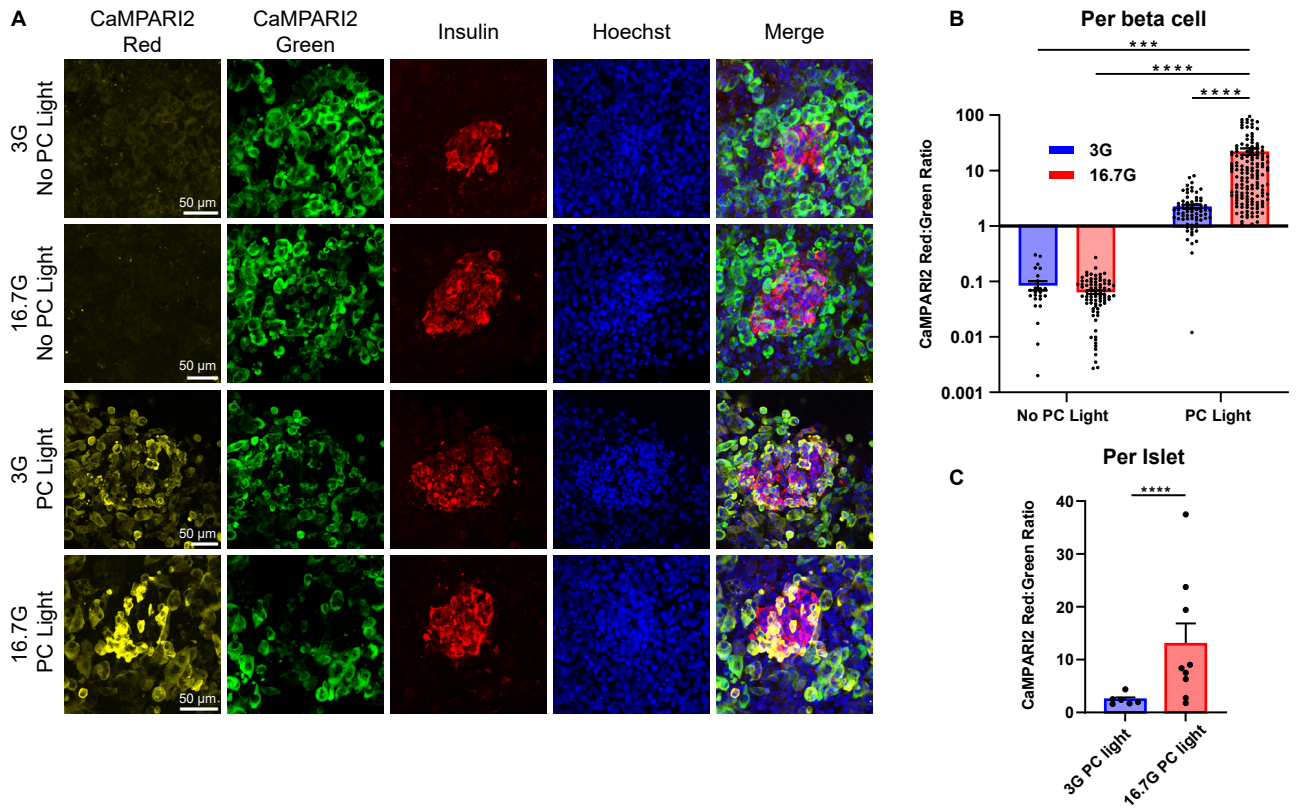

Figure 4: Functional Ca^{2+} Biosensor, CaMPARI2, in Living Human Pancreas Tissue Slices (nPOD 6617 ND). Cells that respond to glucose stimulation experience an influx of calcium ions. An increase in intracellular calcium, along with exposure to ultraviolet (UV) light for photoconversion (PC), irreversibly converts the fluorescence of CaMPARI2 from green to red. **(A)** Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in high or low glucose for ten minutes, followed by exposure or no exposure to PC light. PC light-treated groups were exposed using ten second pulses for five minutes. Tissue slices were then chemically fixed with paraformaldehyde for staining. **(B, C)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 red:green ratio. Only cells that were positive for insulin were quantified (n = 3 slices per group). Data were analyzed per beta cell or per islet. (± SEM, one-way ANOVA with Tukey's post-hoc test, *** $p < 0.001$, **** $p < 0.0001$, outliers removed with ROUT Q = 1%).

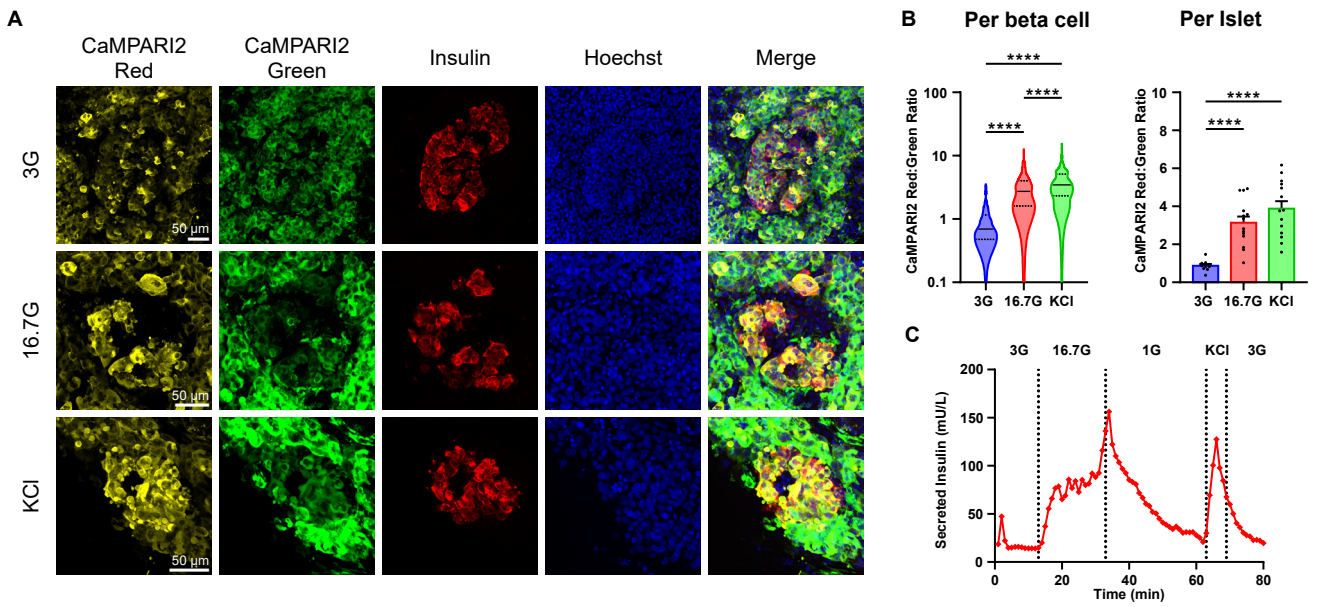

Figure 5: Functional Ca2⁺ Biosensor, CaMPARI2, in Living Human Pancreas Tissue Slices (nPOD 6622 ND Control). (A) Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in 3 mM and 16.7 mM glucose for ten minutes, or 30 mM KCl for one minute. Slices were then exposed to PC light using ten second pulses for five minutes before being washed with PBS. Tissue slices were then chemically fixed with paraformaldehyde for staining. Beta cells displayed strong calcium activity upon high glucose stimulation, resulting in significant photoconversion. **(B)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 Red:Green Ratio. Only cells that were positive for insulin were quantified. (n = 4) slices per group. (± SEM, one-way ANOVA with Tukey's post-hoc test, *** $p < 0.001$, **** $p < 0.0001$, outliers removed with ROUT Q = 1%). **(C)** Perifusion analysis of pancreatic tissue slices from this non-diabetic donor demonstrated robust insulin secretion in response to high glucose stimulation, confirming the preserved glucose responsiveness of beta cells.

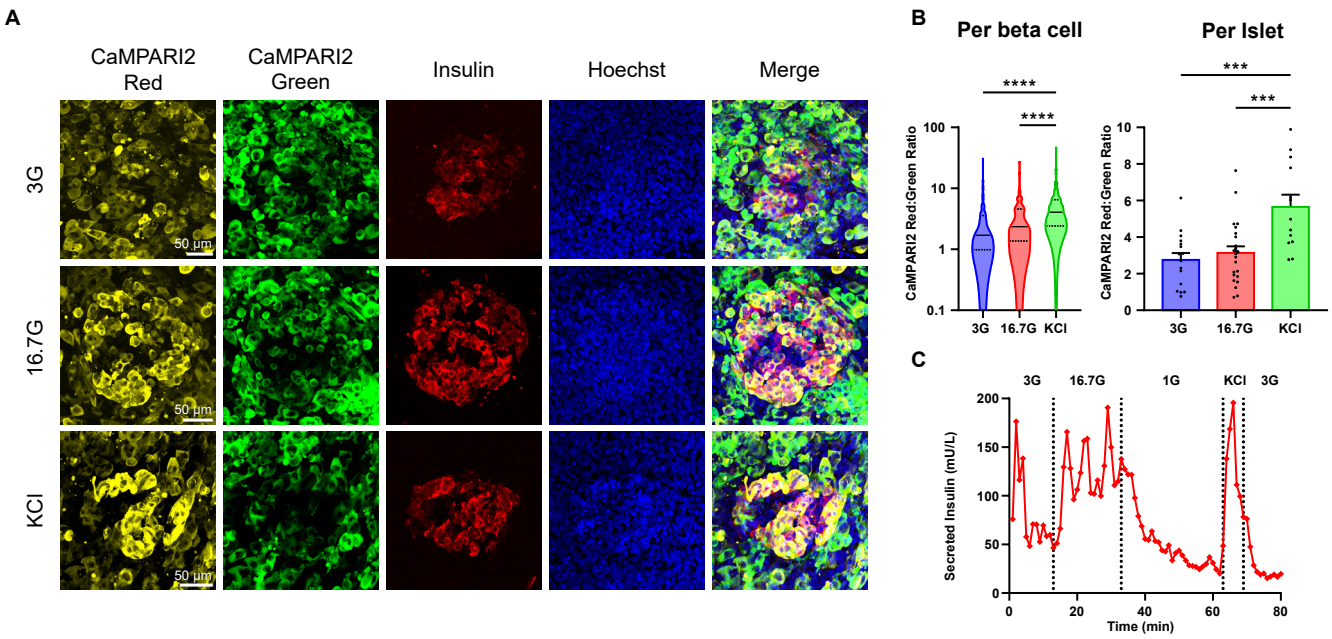

Figure 6: Functional Ca^{2+} Biosensor, CaMPARI2, in T2D nPOD Case 6619. (A) Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in 3 mM and 16.7 mM glucose for ten minutes or 30 mM KCl for one minute. Slices were then exposed to photoconversion (PC) using ten second pulses for five minutes before washing with PBS. Tissue slices were then chemically fixed with paraformaldehyde for staining. **(B)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 Red:Green ratio. We observed a dysregulation in beta cell glucose sensing, particularly elevated calcium influx in low glucose conditions. Only insulin-positive cells were quantified (n = 4 slices per group). Data were analyzed on a per-beta-cell or per-islet basis. (± SEM, one-way ANOVA with Tukey's post-hoc test, *** $p < 0.001$, **** $p < 0.0001$, outliers removed with ROUT Q = 1%). **(C)** The T2D case exhibited an atypical response, with significantly elevated insulin secretion at low glucose levels, aligning with the dysregulated calcium influx observed using CaMPARI2.

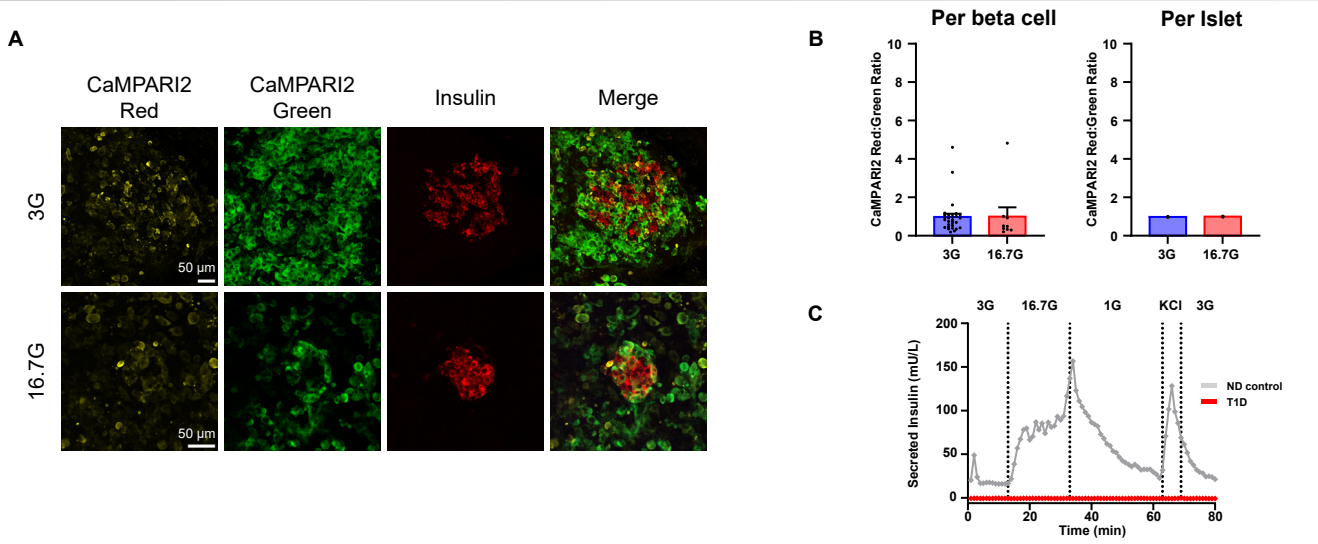

Figure 7: Functional Ca^{2+} Biosensor, CaMPARI2, in T1D nPOD case 6618. (A) Slices transduced with the CaMPARI2 biosensor were incubated at 37°C in 3 mM and 16.7 mM glucose for ten minutes. Slices were then exposed to PC light using ten second pulses for five minutes before washing with PBS. Tissue slices were then chemically fixed with paraformaldehyde for staining. **(B)** Glucose responsiveness in beta cells was quantified by calculating the CaMPARI2 red:green ratio. The residual beta cells were non-functional, showing no response to high glucose stimulation, resulting in no photoconversion of CaMPARI2. Only cells that were positive for insulin were quantified (n = 1 slice per group). Data were analyzed on a per-beta-cell basis. **(C)** Perifusion analysis of pancreatic tissue slices from this T1D case revealed a total absence of insulin release in response to both high glucose and KCl stimulations, confirming the non-functionality of the remaining beta cells (red). Perifusion trace from non-diabetic (ND) case 6622 (grey) provided as a control.