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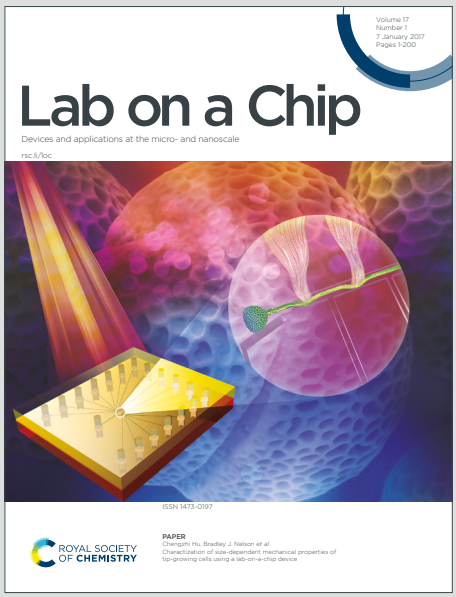

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Microfluidic mimicry of the Golgi-linked N-glycosylation machinery

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Abstract

The complexity of the eukaryotic glycosylation machinery hinders the development of cell-free protein glycosylation since *in vitro* methods struggle to simulate the natural environment of the glycosylation machinery. Microfluidic technologies have the potential to address this limitation due to their ability to control glycosylation parameters, such as enzyme/substrate concentrations and fluxes, in a rapid and precise manner. However, due to the complexity and sensitivity of the numerous components of the glycosylation machinery, very few “glycobiology-on-a-chip” systems have been proposed or reported in the literature. Herein, we describe the design, fabrication and proof-of-concept of a droplet-based microfluidic platform able to mimic N-linked glycan processing along the secretory pathway. Within a single microfluidic device, glycoproteins and glycosylation enzymes are encapsulated and incubated in water-in-oil droplets. Additional glycosylation enzymes are subsequently supplied to these droplets via picoinjection, allowing further glycoprotein processing in a user-defined manner. After system validation, the platform is used to perform two spatiotemporally separated consecutive enzymatic N glycan modifications, mirroring the transition between the endoplasmic reticulum and early Golgi.

1. Introduction

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Protein glycosylation occurs in all domains of life, with N-linked glycosylation being the most abundant type of glycosylation in eukaryotes.[1-3] Such post-translational modifications on the surface of proteins are intimately involved in various processes, including protein folding and its quality control, cell-cell recognition or antibody binding/recognition.[4-9] In eukaryotic cells, protein glycosylation is initiated in the endoplasmic reticulum (ER) by oligosaccharyltransferase (OST) transferring a pre-assembled oligosaccharide, $GlcNAc_2Man_9Glc_3$, onto an asparagine residue of the Asn-X-Ser/Thr amino acid sequon of nascent polypeptides.[10, 11] This N-glycan chain is subsequently used and trimmed in the ER during protein quality control.[5, 6] In the Golgi, further saccharide modifications lead to a variety of different glycan structures covalently linked to proteins.[8, 12, 13] These modifications are dependent on parameters such as substrate and enzyme concentrations, their fluxes through the glycosylation compartments as well as the protein's structure.[12, 14] The influence of protein structure leads to further complexity if multiple glycosylation sites exist. Here, the type and degree of the processing may vary between each site. The resulting site-specific heterogeneity requires the use of bespoke mass spectrometry methods for a detailed characterisation.[12, 14, 15]

Since the glycosylation patterns of glycoproteins serve many functions *in vivo*, they are highly important in therapeutics. Specific forms of glycosylation have been shown to influence and control the effects of therapeutic proteins.[16-19] Indeed, chemoenzymatic approaches have been developed to "glycoengineer" monoclonal antibodies towards more desirable glycan forms, with a focus on achieving homogeneous glycosylation patterns.[20-23] For example, N-linked glycans have been enzymatically hydrolyzed and a drug-glycan conjugate subsequently attached.[24] Nevertheless, substrate specificities of processing enzymes are not universal, and the aforementioned approaches primarily address homogeneous IgG glycosylation for antibody-drug conjugations. However, the presence of heterogeneous glycosylation patterns may be advantageous in certain applications, such as the development of glycoprotein or bioconjugate vaccines.[16, 25, 26] Accordingly, and to better control glycosylation heterogeneity and thus leverage it in therapeutic applications a better understanding of the underlying mechanisms must be developed.

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The traditional approach to controlling glycosylation heterogeneity involves genetic glycoengineering. However, this normally requires extensive cell-line engineering and is limited since such alterations may have unfavourable effects on the cell due to the importance of glycosylation in eukaryotic processes.[16, 25, 26] In parallel, the complexity of the glycosylation machinery makes predictive network engineering demanding and the study of competing glycosylation reactions remains far from trivial. In both genetic and chemoenzymatic glycoengineering, an understanding of the glycosylation machinery and the enzyme kinetics involved is key to understanding and further developing approaches to yield glycoproteins with desired glycosylation patterns. To this end, alternative approaches and techniques able to investigate the processes and effects of glycosylation are required.[16, 26]

In vivo glycosylation parameters, such as protein, saccharide and enzyme concentrations, their availabilities due to competing reactions, and the spatial and temporal separation of reactions between ER and different Golgi cisternae are difficult to control using conventional or “bulk” approaches. Microfluidic technologies, however, allow for the efficient control over reagent (protein and buffer) concentrations, facilitate reaction compartmentalization and enable control over fluxes between user-defined compartments.[27] These features make microfluidic systems particularly interesting as tools to mimic the glycosylation machinery *in vitro*. Indeed, microfluidic technologies have previously been used to synthesize a variety of biomolecules including oligosaccharides and proteins.[28-34]

The earliest example of the use of microfluidics to mimic aspects of the Golgi apparatus was reported by Linhardt and co-workers in 2009.[35] Using a digital microfluidic platform, heparin was enzymatically sulfonated by merging two reactant-containing source droplets. Unfortunately, the low throughput nature and structural complexity of digital microfluidic systems severely limited its application and adoption by others. Additionally, separation of the product from the reaction mixture was achieved by immobilizing the substrate (heparan sulfate) onto streptavidin functionalized magnetic nanoparticles. Although successful, immobilization of the reaction substrate significantly increased the complexity of the workflow. More recently, DeLisa and co-workers presented a continuous flow microfluidic system capable of performing cell-free protein synthesis of superfolder green fluorescent

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protein (sfGFP) and its subsequent glycosylation.[36] In a first module, sfGFP was synthesized from plasmid DNA (encoding the acceptor protein) mixed with a crude yeast cell extract. The reaction product was then delivered to a second module, containing the bacterial OST enzyme, C. jejuni PglB, linked to the surface of the microfluidic channel. Here, a heptasaccharide is transferred from an undecaprenol-pyrophosphate-linked heptasaccharide glycan donor. A final module was then used to isolate the protein product via metal affinity capture. Whilst this study established the feasibility of glycosylation-on-a-chip (transferring an initial glycan to yield a glycoprotein) it should be noted that microfluidic systems have yet to be used to modify the glycosylation profile of glycoproteins. This is in large part due to a number of practical considerations. Of particular importance is the nature of the material used to form the microfluidic device itself. The most common material used to make microfluidic systems is polydimethylsiloxane (PDMS). Unfortunately, PDMS absorbs/adsorbs a wide range of chemical and biological molecules, including proteins.[37-41] This is highly problematic when performing complex reactions or assays since macromolecular concentrations will vary in an uncontrollable manner in both space and time. The extent of biofouling within microfluidic systems is (macro)molecule-specific, posing a challenge when creating surface modification chemistries that can be “universally” employed to prevent biofouling. [37-41] Indeed, since PDMS-based microfluidic systems are applied to a wide range of chemical/biological problems, involving a variety of small molecules and macromolecules, surface modification chemistries must be tailored on case by basis. To address this issue and create a robust and configurable platform for chip-based glycosylations, we describe the fabrication and testing of a PTFE-based microfluidic platform that employs ER and Golgi resident glycoenzymes to alter glycan structures of a model glycoprotein and generate distinct glycosylation patterns. Our microfluidic approach aims to control key parameters associated with the glycosylation machinery, including enzyme and substrate concentrations, temperature and retention times inside defined compartments. Specifically, enzymatic reactions are compartmentalized in water-in-oil droplets that can be incubated at elevated temperatures for defined periods of time. Passive mixing structures ensure proper mixing of the droplet contents, whilst the geometry of the incubation chambers used ensures a stable and uniform flow of droplets through the system. Additionally, by using picoinjectors, additional enzymes or reactants can be added to pre-formed

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droplets and subsequently incubated “on chip”. To demonstrate the efficacy of our platform for chip-based glycosylations, we enzymatically modify N-linked $Man_9GlcNac_2$ glycosylated yeast protein disulfide isomerase (PDI) by the glycosylation enzymes ER mannosidase I (ERMan I) and Golgi mannosidase I (GM I). Glycosylation patterns are then analyzed using tandem mass spectrometry (MS/MS), building on previous work by Hang *et al*.[15] and Mathew *et al*.[14] on the kinetics for these enzymes, we mimic the initial steps of mammalian glycosylation modifications between the ER and Golgi apparatus and use our microfluidic system to study the kinetics of the glycosylation machinery network.

2. Materials & Methods

Recombinant protein and enzyme production and purification

The recombinant glycoprotein protein disulfide isomerase (PDI) and glycoenzymes ERMan I and GM I were constructed, produced and purified as described in previously.[14, 15] Briefly, secreted PDI was transformed into DH10Bac E. coli cells (#10359016, Thermo Fisher Scientific, Switzerland) for bacmid expression. After isolation of bacmid DNA, sf21 cells (#11497013, Thermo Fisher Scientific, Switzerland) were transfected using Cellfectin™ II (Thermo Fisher Scientific, Switzerland) and the baculovirus stock harvested from the supernatant after 72 hours. Baculovirus stocks of N-terminally His_8-tagged, secreted human glycoside hydrolases ERMan I (MAN1B1) and GM I (MAN1A2) were obtained from the glycoenzyme repository.[14, 42] Baculovirus stocks were amplified in sf21 cells to ensure high titers of infectious virus particles.

Protein expression of PDI, ERMan I and GM I was achieved by infecting High-Five™ cells (#B855-02, Thermo Fisher Scientific, Switzerland) through the addition of 1:100 v/v of the respective virus stock to the cell culture. 10 µM of the α-1,2-mannosidase inhibitor kifunensine (Sigma-Aldrich, Switzerland) was added during initial infection of High-Five™ cells with PDI baculovirus to yield a homogeneous PDI glycosylation pattern. Cells were pelleted after 48 hours and then flash-frozen using liquid nitrogen. After cell lysis, with 1% TritonX-100 (Carl Roth, Germany) in phosphate buffered saline solution (PBS, 135 mM NaCl, 2.5 mM KCl, 10 mM Na_2HPO_4, 1.75 mM KH_2PO_4, pH = 7.4), the His10-tagged PDI was purified using Protino® Ni-NTA agarose affinity

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chromatography (Machery-Nagel, Germany) (**Figure S1**).[14] Following purification, PDI was immediately buffer exchanged to PBS (pH = 7.4). Protein concentration was determined using a Nano-Drop Spectrophotometer (Thermo Fisher Scientific, Switzerland). PDI was flash-frozen in liquid nitrogen and stored at -80°C.

The secreted glycosylation enzymes ERMan I and GM I were harvested from the infected High-Five™ supernatant after 72 hours and centrifuged for 10 minutes at 3500 rcf. The supernatant was then filtered using sterile 0.22 µm filters (TPP Techno Plastic Products, Switzerland) and incubated with 2 % v/v Ni-NTA beads (Machery-Nagel, Germany) for 3 hours at 4°C. As with PDI, glycosylation enzymes were purified over their His10-tag using identical Ni-NTA affinity chromatography protocols (**Figure S2**). After purification, ERMan I and GM I were immediately buffer exchanged to their respective activity buffers (**Table S1**). Protein concentrations were determined using a Nano-Drop Spectrophotometer (Thermo Fisher Scientific, Switzerland). The purified glycosylation enzymes were diluted with glycerol (Sigma-Aldrich, Switzerland) (25 % v/v final concentration), flash-frozen in liquid nitrogen and stored at -80°C.

Microfluidic platform design considerations

As described previously, the microfluidic platform comprises four functional modules that allow the encapsulation, mixing, reaction and incubation of substrate glycoprotein and glycosylation enzymes, and the controlled addition of secondary enzymes/reagents after user-defined time periods. Droplets are formed at a flow focusing geometry having a nozzle width of 30 µm and a height of 40 µm. Droplet reaction/incubation times are then defined using a series of constrictions and chambers that redistribute droplets repeatedly as they move along the flow path. This process results in droplet shuffling and provides for control of droplet incubation times without significant incubation time distributions.[43] The developed platform contains two droplet incubation modules containing either 117 or 286 170 µm-high chambers in series. The first incubation module has a total volume of 23 µl, whereas the second incubation module has a volume of 60 µl. Introduction of additional enzymes or reagents into preformed droplets is achieved using a picoinjector that incorporates 1M saltwater electrodes. The height of the picoinjection channel was set equal to the height of the droplet generation module (40 µm) to prevent backflow into the injection

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channel. Features in the rest of the picoinjection module were made to be 80 µm high, so as to limit any backpressure generated in the module.

Microfluidic platform fabrication

Channel patterns were designed using AutoCAD® 2018 software (Autodesk, USA). Master molds were fabricated using a previously described protocol.[44] Briefly, SU-8 photoresist layers (GM1070, Gersteltec, Switzerland) of variable thickness were spin coated on a single silicon wafer (Siegert Wafer, Germany). The layer for inlet channels was 40 µm thick, picoinjection layer 80 µm and the droplet incubation and inlet holes were 170 µm thick. Alignment of different layers was performed using a UV-KUB3 mask aligner (Kloe, France). After master mold fabrication, the entire wafer was exposed to chlorotrimethylsilane (Sigma-Aldrich, Switzerland) vapour for at least 1 hour to aid the removal of PDMS later in the fabrication process. PDMS microfluidic devices were fabricated using standard soft-lithographic techniques. [44, 45] Briefly, this involved casting a PDMS mixture made using a 10:1 w/w ratio of base to curing agent (Elastosil RT 601 A/B, Ameba, Switzerland) onto the patterned silicon wafer and curing for at least 1 hour at 70°C. The cured PDMS was then peeled off the mold, and individual devices were formed by dicing. 0.76 mm diameter inlet and outlet ports were created using a Shaft 20 catheter punch (Syneo, USA). The PDMS replicas were then plasma bonded onto 76 x 26 mm glass slides (Menzel-Glaser, Germany) using a Zepto air plasma (Diener electronic, Germany) and a 120°C post-bake for 4 hours. Immediately after bonding, microfluidic channels were filled and incubated with tridecafluoro-1,1,2,2-tetrahydrooctyl-1-trichlorosilane (abcr, Germany) in HFE 7500 Novec oil (Interelec, Switzerland) for 5 minutes and then post-baked at 120°C for at least 5 hours.

PTFE microfluidic devices were fabricated by adapting a previously described protocols.[46, 47] First, an uncured PDMS mixture (13:1 w/w base to curing agent) containing 40 µl saturated (200 mg/ml in EtOH) Pluronic F127 (Sigma-Aldrich, Switzerland) per 10 grams PDMS mixture was cast onto the SU-8 master mold. The PDMS-Pluronic mixture was then cured for at least 1 hour at 70°C and then removed from the master mold. The cured PDMS replica was then heated for 2 minutes in a microwave oven at 700 W. Subsequently, a double-negative PDMS mold was fabricated by casting uncured PDMS (13:1 w/w base to curing agent) containing 40 µl

saturated Pluronic™ F127 per 10 grams PDMS onto the initial Pluronic-containing PDMS block. This double-negative PDMS mold was cured for 90 minutes, cooled to room temperature over 30 minutes and subsequently removed from the initial PDMS negative mold. Embedding Pluronic™ F127 within the PDMS passivates the PDMS surfaces and reduces polymer movement from one PDMS mold into the double-negative mold during curing. This improves the subsequent separation of the two PDMS blocks. Such a strategy allowed the fabrication of microfluidic channel structures with low width to height ratios. Inlet and outlet ports were created in the double-negative PDMS mold using a Shaft 20 catheter punch (Syneo, USA). Metal pins were inserted into these holes and THV 500GZ PTFE pellets (3M, Germany) melted onto the double-negative PDMS mold overnight in a Vacucenter VC50 vacuum oven (SalvisLab, Switzerland) at 200°C. Next, the molten PTFE block was cooled to room temperature and the metal pins removed using household pliers. The PTFE block was then placed face down onto a flat THV 500GZ sheet that was spin coated from a 5% THV 221GZ PTFE solution in acetone (3M, Germany). The two parts were pressed together with minimal pressure using a custom-made bonding device based on Ren *et al*.[47] but with metal instead of glass plates. The complete PTFE device was thermally bonded at 115°C for at least 2 hours. After bonding, hollow metal connections (Chuang Mei Wei Technology, China) having an outer diameter of 0.76 mm were connected to 0.56 mm ID PTFE tubing (Rotima, Switzerland) and inserted into the fully bonded PTFE device.

Microfluidic platform characterization

Operational testing of the microfluidic system was performed using an Eclipse Ti-E inverted microscope (Nikon, Switzerland) equipped with a Plan Fluor 4x/0.13 objective (Nikon, Switzerland) or a Plan Fluor 10x/0.3 objective (Nikon, Switzerland) and a MotionPro Y5 high-speed camera (IDT Vision, USA). A Dino-Lite digital microscope (AnMo Electronics, Taiwain) was used to monitor droplets during enzymatic assays.

Enzymatic reactions

All enzymatic assays were performed at 42°C by placing the entire microfluidic device on a hot plate. Prior to use, the respective glycosylation enzymes ERMan I and/or GM I were diluted to 22.5 µg/ml in the respective enzyme activity buffer (**Table S1**) and

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2.25% v/v glycerol. PDI (1 mg/ml) and ERMan I (22.5 µg/ml) and/or GM I (22.5 µg/ml) were taken up in 0.56 mm ID PTFE tubing (Rotima AG, Switzerland) connected to 1 mL Hamilton glass syringes (Sigma–Aldrich, Switzerland) containing water (MicroPure™ UV/UF 0.2 µm, Thermo Scientific, Germany) as a supporting fluid. An air bubble between the sample and the supporting fluid in the PTFE tubing prevented contamination and dilution of the sample by the supporting fluid whilst allowing for complete sample consumption. For experiments involving only one glycosylation enzyme, an equivalent blank buffer solution was prepared accordingly.

The oil phase, consisting of 1:1 mixture of droplet generation oil (Bio-Rad Laboratories, Switzerland) and HFE 7500 Novec oil, was taken up in a 1 mL Hamilton glass syringe. Precision neMESYS syringe pumps (CETONI, Germany) were used to move all fluids and provide for a stable flow rate of 0.8 µl/min for each fluid. This resulted in an overall flowrate through the incubation chambers of 2.4 µl/min prior picoinjection and 3.2 µl/min post picoinjection. For picoinjection, a voltage of 60 V at 1 kHz was applied to the saltwater electrodes. Droplets were collected in 40 µl 100 % trichloroacetic acid (500 µg in 227 µl H_2O, Sigma-Aldrich, Switzerland) and 60 µl PBS. The aqueous phase was isolated from the oil phase using 1H,1H,2H,2H-perfluorooctanol (Apollo Scientific Ltd, UK). The recovered protein solution was precipitated in 15 % v/v trichloroacetic acid for 10 minutes and pelleted for 5 minutes at 20'000 rcf and 4°C. The resulting protein pellet was washed 3 times with acetone, air dried and stored at 20°C as previously described.[14]

Mass spectrometry measurements and glycoform quantification

For MS analysis, precipitated protein pellets were resuspended in 400 µl of 8 M urea (Sigma-Aldrich, Switzerland). Sample was then processed according to a previously described protocol.[14] Briefly, proteins were first reduced in 50 mM dithiothreitol (Sigma-Aldrich, Switzerland) followed by alkylation in 130 mM iodoacetic acid (Sigma-Aldrich, Switzerland) and 50 mM ammonium bicarbonate (Sigma-Aldrich, Switzerland) for 30 minutes at 37°C to facilitate trypsin digestion using a 1:80 trypsin (Promega, Switzerland) to PDI weight ratio (overnight at 37°C). Following tryptic digestion, the resulting peptides were desalted with 0.6 µL of Zip-Tip C18 resin (Milipore, Ireland) and dried until use. MS/MS measurements were performed using a calibrated Q Exactive mass spectrometer (Thermo Fisher Scientific, Switzerland) coupled to a

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Acquity UPLC M-Class system (Waters AG, Switzerland) with a PV-500 Picoview nanospray source (Sciex, USA).

MS/MS data were analyzed using Xcalibur 4.0 software (Thermo Fisher Sceintific, Switzerland) as described previously.[14, 15] Spectral peak areas were defined manually and integrated. For simplicity, only the relative amounts of glycoforms on the glycosylation site 2 peptide are investigated in the current study. Additional information regarding the investigated glycosylation site 2 peptide is provided in **Table S2** and in Mathew *et al.*[14]

Assessment of protein adsorption "on-chip"

For experiments in PDMS and Teflon-based microfluidic devices PDI was expressed in High-Five™ cells using a baculovirus expression system described above. For the adsorption experiments PDI was not co-expressed with the α-1,2-mannosidase inhibitor kifunensine (Sigma-Aldrich, Switzerland).

Recombinant PDI and commercially available bovine serum albumin (Sigma-Aldrich, Switzerland) were fluorescently labelled by incubating 15.9 µM PDI or BSA in PBS (pH = 7.4) with 100 molar equivalents of NHS modified Atto 488 in DMSO (ATTO-TEC, Germany) at 37°C for 4 hours followed by four buffer exchanges to PBS (pH = 7.4).

For protein adsorption experiments, 50 µm wide and 40 µm high microfluidic channels were initially flushed with PBS (pH 7.4). This was then replaced by a previously prepared fluorescent protein solutions and incubated for five minutes. Subsequently, channels were flushed with 10 µl PBS (pH 7.4). Residual fluorescence originating from microchannels after flushing was quantified and compared to the pre-incubation background measurement. Fluorescence detection was performed using an Eclipse Ti-E inverted microscope (Nikon, Switzerland). Fluorescence emission was collected using a Plan Fluor 10x/0.3 objective (Nikon, Switzerland), filtered through 469/35 excitation and 525/39 emission filters (IDEX Health & Science, USA). Fluorescence emission was detected using an ORCA-flash 4.0 CMOS camera (Hamamatsu, Solothurn, Switzerland). µManager 1.4 software was used to control and automate fluorescence collection, while ImageJ software (National Institutes of Health, USA) was used for image processing and analysis.

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3. Results

Microfluidic substrate

As discussed previously, the microfluidic platform aims to enable spatiotemporal control over protein concentrations by compartmentalizing enzymatic reactions in droplets and employing picoinjection to add additional enzymes, reactants, substrates, or buffers. A basic consideration in this regard is the choice of substrate material. Despite the widespread adoption of PDMS by the microfluidics community, biofouling of microfluidic channel surfaces will severely compromise the study of biochemical reaction networks. As expected, the use of PDMS-based microfluidic devices in preliminary experiments was characterized by the depletion (adsorption/absorption) of enzyme prior to droplet formation at the flow focusing geometry. Such biofouling impaired downstream enzymatic reactions in droplets (**Figure S3B-D**). To assess the likelihood and/or magnitude of biofouling, 15.9 μM fluorescently labelled protein disulfide isomerase (PDI) and bovine serum albumin (BSA) solutions were separately incubated in 50 μm x 40 μm cross-section microfluidic channels within PDMS and PTFE substrates for 5 minutes. Subsequently, channels were flushed with 10 μl PBS and residual fluorescence measured. As shown in **Figure 1A,C**, Atto-488-labelled BSA remains on or in the microfluidic substrate after washing for PDMS (top) but not for PTFE (bottom). In contrast, only a slight increase in fluorescence after incubation with Atto-488-labelled PDI (and washing) was seen for the PDMS substrate (top), and even less for the PTFE substrate, indicating reduced biofouling (**Figure 1B,C**).

Microfluidic platform

Inside a microfluidic chip, substrate glycoprotein and glycosylation enzymes were co-encapsulated and subsequently incubated “on-chip” for an extended period of time of up to 30 minutes. As glycan hydrolysis is rarely a terminal reaction and further glycan processing by the same enzyme may occur over time, “off-chip” storage of reaction intermediates was limited. The addition of subsequent reaction mixtures was therefore implemented within the same microfluidic chip. A schematic of the entire microfluidic platform is presented in **Figure 2**. It comprises four adaptable modules. In the first, droplets are generated at a flow focusing geometry with substrate and enzyme being co-encapsulated and passively mixed by chaotic advection within a winding channel

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section.[46, 48] Each reagent flux can be independently controlled by the user to regulate droplet payloads. The second module integrates a series of incubation chambers that ensures a stable flow of droplets over an extended incubation period. The average residence time of droplets in this incubation module is controlled by the inlet flow rates and the number of incubation chambers.[43] In the current study, the total residence time of droplets in the incubation module was approximately 10 minutes. The third module contains a picoinjector and allows the controlled addition of small volumes of additional enzyme, substrate, or buffer to the droplets emerging from the first incubation module.[49, 50] Picoinjection works by flowing droplets past a channel containing a pressurized reagent stream. If a droplet is enveloped by a surfactant layer, this fluid stream will normally not enter the droplet. However, application of an electric field can be used to destabilize and breach the surfactant layer, allowing the reagent stream to enter the droplet over a short period of time. The process is highly robust and allows controlled addition of femtolitre–picolitre volumes at kilohertz rates. To reduce the probability of droplet fusion during picoinjection, a grounding electrode that acts as a shielding electrode is employed.[50, 51] The final module has a similar structure to the second module and provides for the controllable incubation of droplets for periods up to 20 minutes after picoinjection.

Biosynthetic system

To prove principle, we focused our attention on glycan processing enzymes of the ER and early Golgi glycosylation machinery. While ERMan I hydrolyses one terminal Mannose on the N-linked glycan $Man_9GlcNAc_2$ down to $Man_8GlcNac_2$, GM I can hydrolyse the glycan further to yield $Man_5GlcNAc_2$,[14, 52] as shown in **Figure 3**. Due to the fact that the ER and Golgi are separated in living systems, a combined mode of action is possible where ERMan I cleaves off a first mannose in the ER, with GM I subsequently trimming the glycan further within the Golgi apparatus. Additionally, it should be noted that prolonged incubation of $Man_8GlcNAc_2$ with ERMan I can lead to further mannose trimming *in vitro*.[14]

First, the glycoprotein substrate (PDI) was expressed in presence of the α-1,2-mannosidase inhibitor (kifunensine) to yield homogeneously glycosylated PDI bearing the N-linked $Man_9GlcNAc_2$ structure. For simplicity, we focused our analysis on glycosylation site 2 of PDI. Enzymatic assays were performed entirely within the

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microfluidic device and followed by droplet collection and protein precipitation off-chip. Glycosylation patterns were then analysed by tandem mass spectrometry (MS/MS).

Microfluidic mimicking of the glycosylation machinery

As described, we first validated our microfluidic platform by performing the enzymatic reactions shown in **Figure 3** within a first-generation microfluidic device shown in **Figure S4**. First, PDI was co-encapsulated with either ERMan I or GM I at the flow focusing geometry, while a blank buffer solution was used in the picoinjector leading to a total incubation time of 18 minutes. The resulting glycosylation patterns for glycosylation site 2 of PDI are shown in **Figure 4A**. As expected, incubating PDI with ERMan I inside droplets yielded $Man_8GlcNAc_2$ as the dominant N-linked glycan. The incubation of PDI and GM I yielded a more heterogeneous glycosylation pattern, with a slow trimming of the first mannose, but accelerated trimming of further mannoses. As a control and to exclude any effects that picoinjection may have on enzyme reactivity, droplets containing the PDI substrate without glycosylation enzymes were picoinjected with ERMan I and GM I. The resulting glycosylation pattern shown in **Figure 4B (left)** indicates that picoinjection has a negligible effect on enzyme activity. The slightly reduced processing of $Man_9GlcNAc_2$ by ERMan I can be explained by the shorter incubation time of 8 minutes due to the absence of enzyme in the droplets during the first incubation module. We subsequently used the final microfluidic device shown in **Figure 2** to mimic the early steps of N-linked glycan processing by the glycosylation machinery. In this proof-of-concept, we investigated the sequential actions of ERMan I and the Golgi resident GM I. Compartmentalization inside droplets ensured their spatial and temporal separation, with droplets containing the PDI substrate and ERMan I being initially incubated on-chip for 10 minutes. The subsequent addition of GM I being achieved by picoinjection was followed by the second incubation for 20 minutes. As shown in **Figure 4C**, the combined and sequential action of the two glycosylation enzymes greatly enhances the hydrolysis of $Man_9GlcNAc_2$, with $Man_8GlcNAc_2$ and $Man_5GlcNAc_2$ being the dominant hydrolysis products. These findings suggest increased reaction kinetics of GM I for the conversion of $Man_7GlcNAc_2$ to $Man_6GlcNAc_2$ and $Man_5GlcNAc_2$, which is in good agreement with recent investigations into the reaction kinetics of the employed glycosylation enzymes.[14]

4. Discussion

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In the current study, we have presented a droplet-based microfluidic platform able to mimic the transition between the ER and early Golgi by controlling the spatiotemporal separation of enzymatic reactions with ER and Golgi resident mannosidases. When mimicking mammalian glycosylation, it is important to consider that glycosylation enzymes are sensitive to environmental conditions and produced at relatively low concentrations in cell culture. This dictates that enzymes should only minimally interact with substrate material of the microfluidic device to avoid their depletion. Any biofouling alters reaction composition in the microfluidic system, which in turn alters or prevents enzymatic reactions. The use of droplet-based microfluidics addresses this issue in part by encapsulating enzymatic reactions and minimizing the interaction between enzyme and microfluidic channel surfaces. However, prior to droplet formation both enzyme and substrate contact the microfluidic channel walls. Such residual biofouling can have severe effects, especially or when processing sensitive enzymes or working with low enzyme concentrations. Accordingly, PDMS is poorly suited for use in such investigations due to its propensity to biofoul. Instead, we fabricated of PTFE-based microfluidic devices, provides access to a far wider range of glycosylation reactions. Although the developed microfluidic platform is able to mimic aspects of the early Golgi-linked glycosylation machinery, the capacity to control molecular fluxes spatially and temporally, means that it can be easily adapted to investigate other parts of the glycosylation machinery and more complex enzymatic networks. For example, the current platform could be extended to incorporate further enzymatic reactions such as those involving glycosyl transferases. Picoinjection could be used to supply reaction droplets with glycosyl transferase (GnT I) and its corresponding saccharide substrate (UDP-GlcNAc). Here, the attachment of a first GlcNAc to the N-linked glycan is crucial in forming hybrid and complex N-linked glycans.[53-55] Picoinjection of kifunensine at varying concentrations could then be used to inhibit glycan trimming by preceding α-1,2-mannosidases. Additionally, enzymes could be removed from droplets using the enzyme's Strep II Tag, available in the employed expression system of the glycoenzyme repository.[42] Here, magnetic nanoparticles functionalized to bind the Strep II Tag could be picoinjected into the droplets, with enzyme-nanoparticle complexes subsequently being removed through the use of magnetic fields and asymmetric droplet splitting as described by Choi and co-workers.[56] Alternatively,

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enzyme nanoparticle conjugations could be used to perform glycosylation reactions. From an analytical standpoint, further insight would be gained through expanding the analyzed glycosylation patterns to other glycosylation sites. In the current study, we focused our analysis on glycosylation site 2 of PDI. Extension to the other four glycosylation sites of PDI would likely yield additional information about surface-specific enzyme-protein interactions.[12, 14] Finally, our work can also be seen in light of recent advances where microfluidic systems have been used to synthesize glycoproteins.[36] For example, by combining our work with such advances, one could possibly synthesize glycoproteins *in vitro* and subsequently study their processing within an integrated microfluidic platform. Finally, enzyme-functionalized magnetic nanoparticles could be co-encapsulated with the reactants in our microfluidic platform. Using asymmetric droplet splitting, one could then also regenerate the precious enzymes for further experiments.

Data Availability: The data underlying this article will be shared on reasonable request to the corresponding author.

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Abbreviations: Asn, asparagine; BSA, bovine serum albumin; ER, endoplasmic reticulum; ERMan I, ER mannosidase I; Glc, glucose; GlcNAc, N-acetylglucosamine; GM I, Golgi mannosidase I; Man, mannose; MS, mass spectrometry; MS/MS, tandem mass spectrometry; OST, oligosaccharyltransferase; PBS, Phosphate buffered saline; PDI, protein disulfide isomerase; PDMS, polydimethylsiloxane PTFE,

polytetrafluorethylene; Ser, serine; sfGFP, superfolder green fluorescent protein; Thr, threonine.

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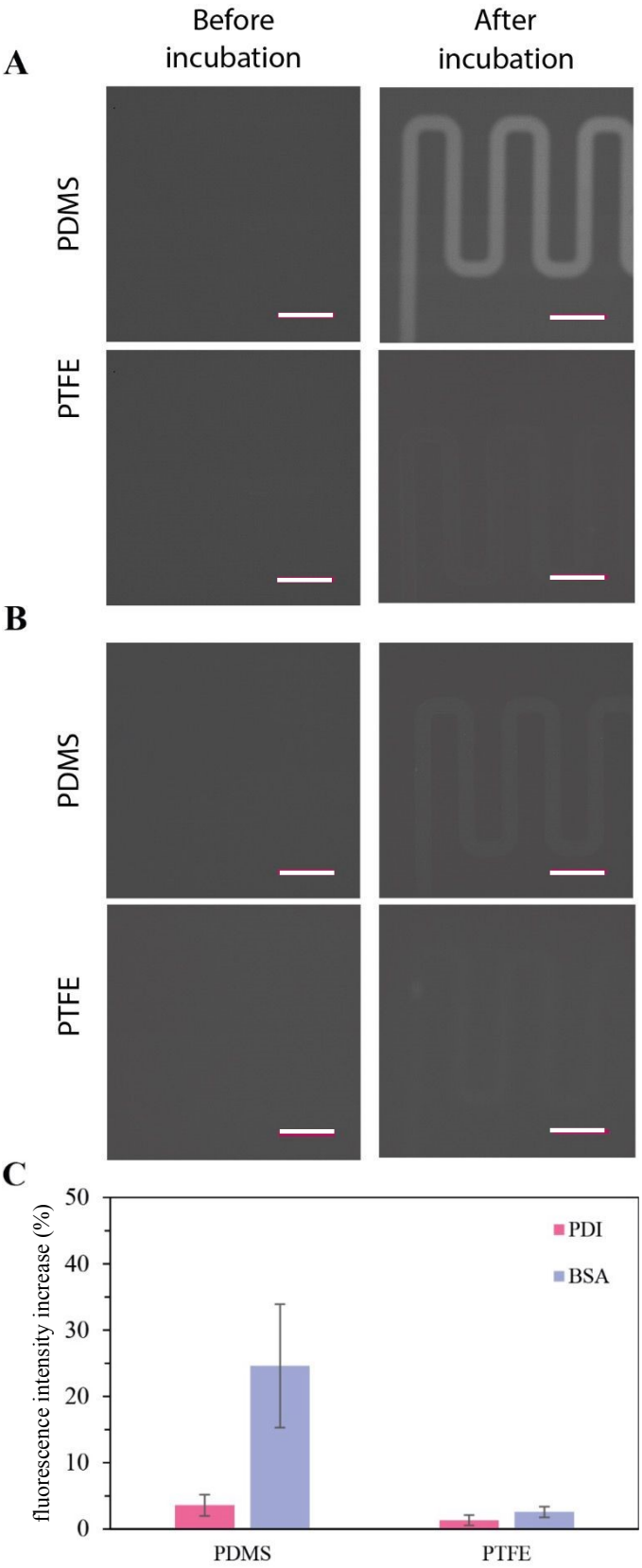

Figure 1: Adsorption/absorption of Atto 488-labelled BSA and Atto 488-labelled PDI to the walls of a 50 µm x 40 µm cross-section microfluidic channel. (A) PDMS and

PTFE channels before and after incubation with Atto-488-labelled BSA and subsequent flushing with PBS. All four images are shown with the same contrast settings. (B) PDMS and PTFE channels before and after incubation with Atto-488 labelled PDI and subsequent flushing with PBS. All four images are shown with the same contrast settings. Scale bars are 200 microns. (C) Percentage increase in time-integrated fluorescence intensity reporting the adsorption/absorption of Atto 488-labelled BSA and Atto 488-labelled PDI to the walls of a 50 μm x 40 μm cross-section microfluidic channel. Fluorescence originating from PDMS increased significantly after incubation with Atto-labelled BSA, with a more moderate increase observed for PDI. In comparison, a negligible increase in fluorescence is observed when PDMS is replaced by PTFE. Error bars represent one standard deviation for triplicate measurements.

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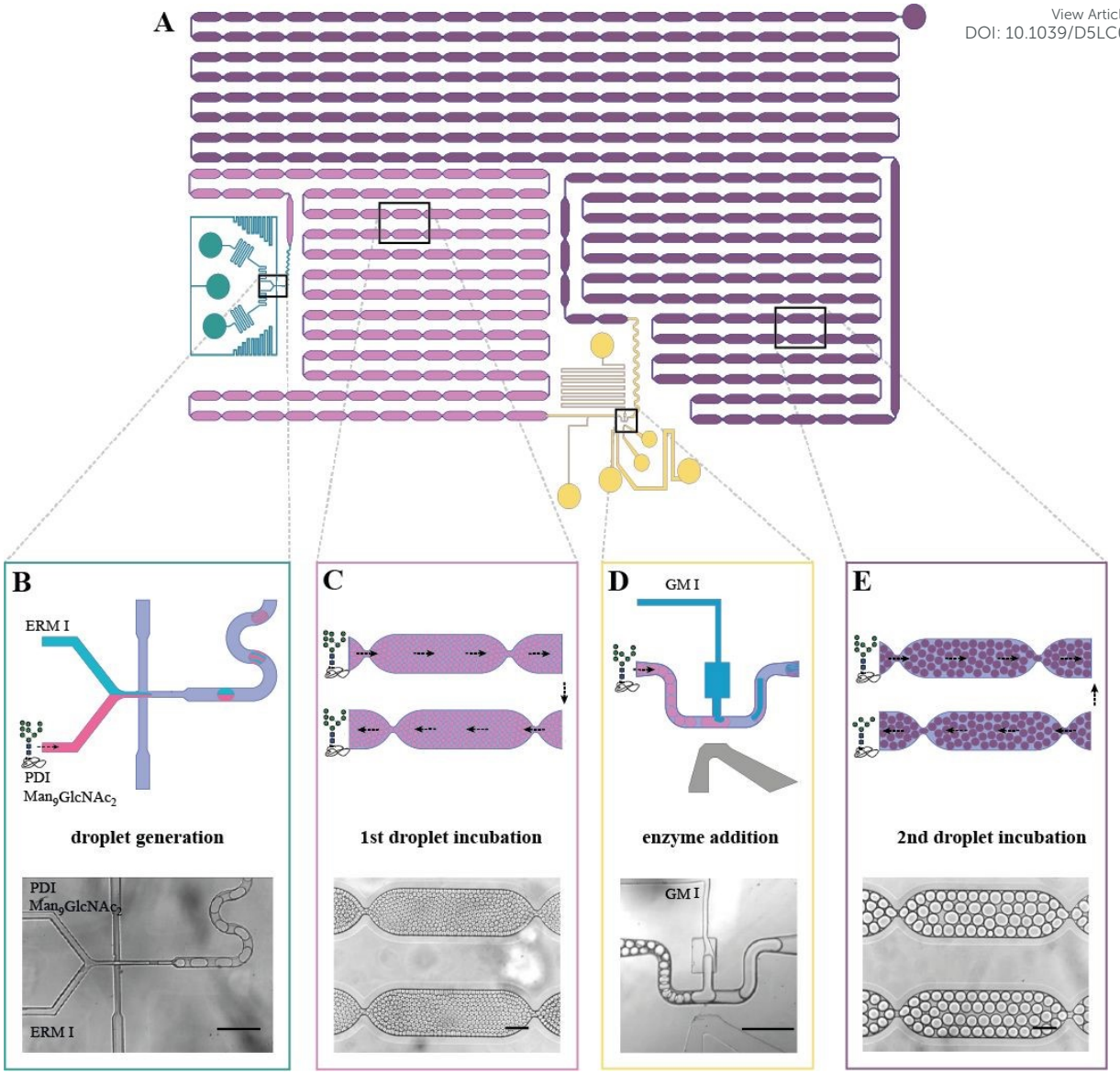

Figure 2: A microfluidic platform for performing enzymatic glycosylation reactions on protein-linked glycans. (A) Schematic of the entire PTFE microfluidic device integrating four droplet processing. (B) Schematic and brightfield image of the flow focusing geometry used to form droplets and coencapsulate the glycoprotein substrate and glycosylation enzyme. (C) Schematic and brightfield image of two chambers within the first incubation module. Incubation times of up to 10 minutes can be realised using the structure shown. (D) Schematic and brightfield image of the picoinjector used to deliver additional enzyme, substrate, and buffer into pre-formed droplets; (E) Schematic and brightfield image of two chambers in the second incubation module. Incubation times of up to 20 minutes can be realised using the structure shown. Scale bars are 300 microns.

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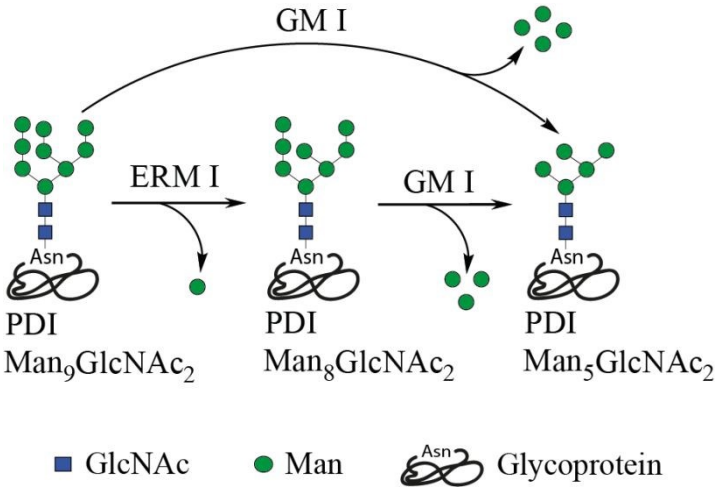

Figure 3: Enzymatic reactions of the biosynthetic system studied. The ER-resident mannosidase ERMan I preferentially hydrolyses a terminal mannose on the substrate $Man_9GlcNAc_2$ yielding $Man_8GlcNAc_2$. In contrast, the Golgi mannosidase I (GM I) is able to hydrolyse $Man_9GlcNAc_2$ to $Man_5GlcNAc_2$, but with a slow hydrolysis of the first terminal mannose.

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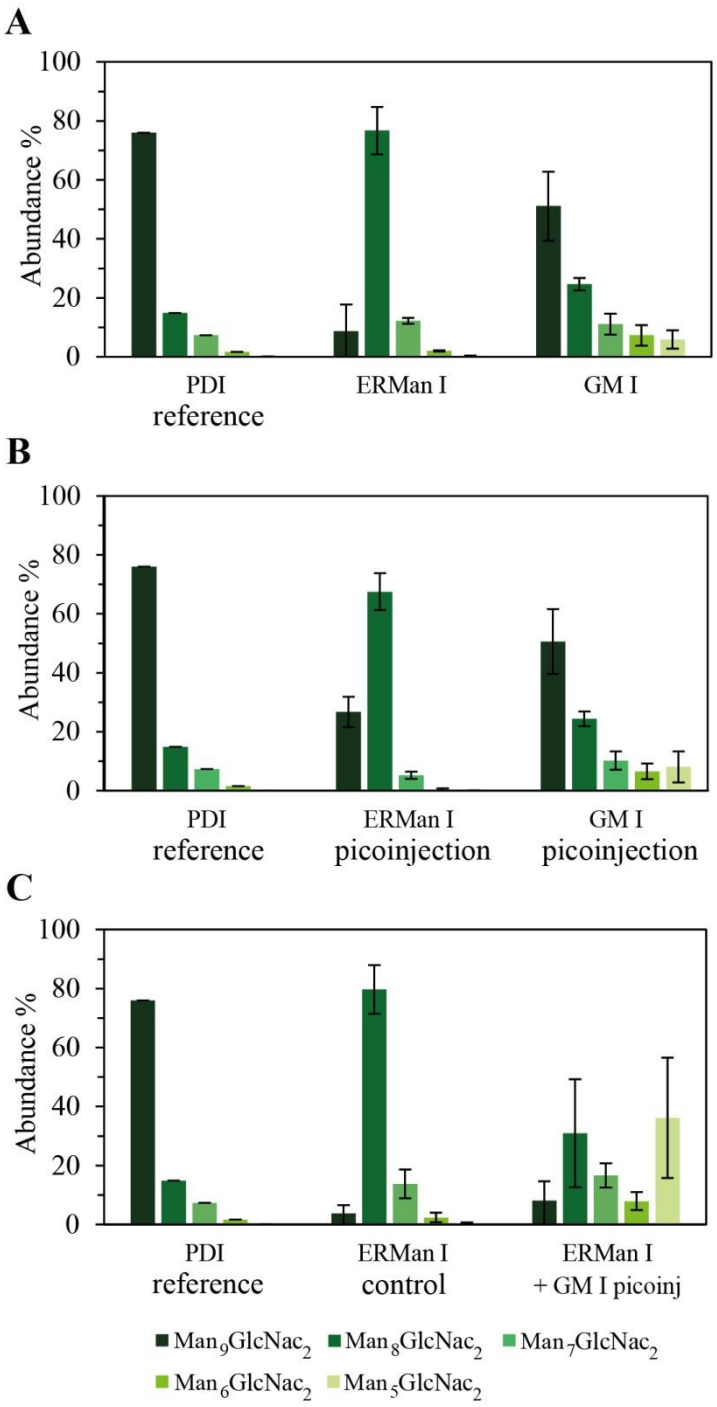

Figure 4: Protein glycosylation patterns after glycosylation reactions on-chip. (A) Co-encapsulation of the glycoprotein substrate PDI-$Man_9GlcNAc_2$ (PDI reference) with glycosylation enzymes ERMan I and GM I. (B) Picoinjection of ERMan I or GM I into droplets containing PDI instead of initial co-encapsulation of enzyme with PDI. (C) Mimicking *in vitro* the *in vivo* glycosylation pathways by co-encapsulation and incubation of PDI with ERMan I followed by picoinjection of GM I and subsequent incubation. As a negative control the glycosylation of PDI after co-encapsulation and

incubation of PDI with ERMan I and subsequent picoinjection of blank enzyme buffer is shown.

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Microfluidic mimicry of the Golgi-linked N-glycosylation machinery

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Data availability statement

Data supporting this article have been included as part of the ESI