Cellular Senescence and Senescence-Associated Secretory Phenotype Drive Multipotent Cardiac Stem/Progenitor Cell Dysfunction in Human Diabetic Cardiomyopathy Independently of Age

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Abstract

Background: Aging and Diabetes Mellitus (DM) independently and additively increase cardiovascular risk and a pathophysiological basis for this epidemiological link is currently the subject of intense investigations. Both aging and DM affect the biology and regenerative potential of tissue specific cardiac adult stem/progenitor cells (CSCs). In aged subjects over half of the CSCs are senescent with a senescence-associated secretory phenotype (SASP) that renders otherwise healthy CSCs to senescence, impairing their proliferative and differentiation potential. Although a link to organismal aging is clear, cells can undergo senescence, regardless of age. Yet, it is unclear whether the SASP is induced by Diabetes per se in CSCs and whether targeting senescent cells within the diabetic CSC compartment rescues their regenerative biology.

Methods: In order to investigate the effects of DM on CSC senescence and to try separate those of aging from Diabetes on senescence events, we obtained peri-infarct/border zone biopsies from non-aged patients (50-64 years old) with DM type 2 (T2DM) and non-diabetic (NDM) patients with post-infarct cardiomyopathy undergoing surgical coronary revascularization.

Results: Ischemic injury in DM is associated with a higher ROS production, as revealed by the exacerbated expression of 8-OH-deoxyguanosine, nitrotyrosine, and 4-hydroxynonenal targeting both cardiomyocytes as well as CSCs. The latter associated with an increased number of senescent and dysfunctional T2DM-hCSCs (isolated from atrial samples of T2DM and NDM patients with ischemic cardiomyopathy) identified by increased p16INK4a positive cells, reduced telomerase activity and telomere length, reduced proliferation, clonogenesis/spherogenesis and myogenic differentiation when compared to NDM-hCSCs in vitro. Importantly, T2DM-hCSCs show a defined SASP, as demonstrated by the increased secretion of MMP-3, PAI1, IL-6, IL-8, IL-1β and GM-CSF. A combination of two senolytics, Dasatinib and Quercetin, clears senescent T2DM-hCSCs in vitro restoring expansion and myogenic differentiation capacities of the diabetic hCSC pool.

Conclusions: DM hampers human CSC biology, inducing a variety of hallmarks of senescence in non-aged subjects that contribute to the deficit of their regenerative potential. Clearance of senescent cells by senolytics abrogates the SASP and restores a fully proliferative- and differentiation- competent hCSC pool in T2DM.

Background

Aging is a major risk factor for the occurrence of several diseases, including stroke, myocardial infarction, heart failure, neurodegenerative disease, and several cancers. Therefore, the global rise in life expectancy will lead to a dramatic increase in age related diseases in the coming decades. In this scenario, cardiovascular diseases including atherosclerosis and heart failure (HF) increase exponentially with age, whereby HF is considered an epidemic because of a growing and ageing population. Type 2 diabetes mellitus (T2DM) is closely bound to aging and is a powerful independent risk factor for
cardiovascular diseases such as atherosclerosis and heart failure [1]. This epidemiological connection between aging, T2DM, and cardiovascular diseases postulates that there might be a pathophysiological link.

Aging has been associated with systemic inflammation and oxidative stress [2], which can be both a cause as well as a consequence of Diabetes [3]. Cellular senescence can be defined as a permanent arrest of cellular growth and is a key feature of aging [4, 5]. Cell senescence is also a cause and a consequence of Diabetes and plays an important role in its cardiovascular complications. Although senescent cells are classically reported as cells that irreversibly cease proliferation, they have the capacity to produce and secrete soluble factors that can influence neighboring cells and tissues [6, 7]. This feature of senescent cells to secrete these soluble factors has taken the name of senescence-associated secretory phenotype (SASP) [8]. Since chronic inflammation is an important pathophysiological factor of both aging and diabetes, the SASP has been pointed as the pathophysiological link between aging and diabetes in cardiovascular diseases [9].

Tissue-specific adult stem cell senescence has emerged as an attractive theory for the decline in mammalian tissue and organ function during aging [10]. The mammalian heart, including the human, harbors a tissue specific cardiac stem/progenitor cell (CSC) compartment [11–15] that undergoes senescence with age, which dictates a progressive and permanent dysfunction of more than half of these endogenous cells by 75 years of age [5, 10, 16, 17]. The senescent CSCs exhibit a SASP that can negatively impact surrounding cells, causing otherwise healthy and cycling-competent CSCs to lose proliferative capacity and switch to a senescent phenotype. Nevertheless, even at the oldest age, it is still possible to retrieve a healthy, cycling-competent CSC fraction with an increased regenerative and reparative capacity [5, 10, 16]. Accordingly, experimental selective ablation of senescent CSCs either genetically or by a combination of senolytic drugs, fosters the expansion and functional regenerative recovery of the healthy aged CSCs [5, 10, 18].

Several reports have shown that Diabetes impairs the in vitro proliferative and differentiation potential of adult CSCs, further worsening their senescence phenotype even when compared with CSCs from non-diabetic ischemic patients [19–21]. Changes in chromatin conformation underlie the impaired proliferation, differentiation, and senescent behavior of diabetic CSC [22]. Yet, it is unclear whether the SASP is induced by Diabetes per se in CSCs and whether targeting senescent cells with the diabetic CSC compartment rescues CSC proliferation and differentiation defect in T2DM. On this premise here we show that the myocardium of T2DM patients, in a very narrow age window of 55–64 years, undergoing cardiac revascularization for ischemic heart disease, is characterized by an increased oxidative stress that affects the CSC compartment with an increased senescent phenotype when compared to myocardium and CSCs of age and sex matched NON-Diabetic Mellitus (NDM) patients. Importantly, Diabetic-Senescent CSCs have a SASP and senolytic treatment abrogate the senescent diabetic Diabetic-Senescent CSCs uncaging healthy CSCs to proliferate and differentiate normally.

Methods
Patients Cohort and Samples

Human myocardial samples were obtained from Type 2 diabetes mellitus (T2DM) and NON-Diabetic Mellitus (NDM) patients with post-infarct cardiomyopathy which undergoing surgical coronary revascularization. The samples were collected only when removal of tissue was required for surgical reasons. Collection of human tissues samples was approved by the local ethics committee at the University of Campania "L. Vanvitelli" of Naples. Before cardiac surgery, written informed consent was obtained from every patient. All patient data were kept anonymous. We included 10 T2DM and 6 NDM patients from which peri-infarct/border zone biopsies were obtained (Table 1). Freshly excised samples were formalin fixed for immunohistochemistry analysis as described below. Furthermore, additional 6 T2DM and 6 NDM patients with similar characteristics were included (Table 1), from which atrial samples were obtained and processed for cell harvesting as below described.

Human CSCs Isolation

Human samples obtained from T2DM and NDM patients were stored in saline on ice until ready to process (~1hr). All steps were performed at 4°C unless stated otherwise. Briefly, cardiac tissue was minced then digested with collagenase II (0.3mg/ml; Worthington Laboratories) in Dulbecco's Modified Eagle's Medium (DMEM; Sigma-Aldrich) at 37°C in a series of sequential digestions for 3 minutes each. Enzymatically released cells were filtered through a 40µm cell strainer (Becton Dickinson, BD) to eliminate the cardiomyocyte population and collected in enzyme quenching media (DMEM + 10% FBS). The isolated cardiac cells were collected by centrifugation at 400g for 10 min, resuspended in incubation media (PBS, 0.5% BSA, 2 mM EDTA). For the isolation of c-kit<sup>pos</sup>CD45<sup>neg</sup>CD31<sup>neg</sup> CSCs the MACS technology was used (Miltenyi Biotec). First, cardiomyocyte-depleted cardiac cells were negatively sorted for CD45<sup>pos</sup> and CD31<sup>pos</sup> cells by immunolabelling with anti-human CD45 and CD31 magnetic immunobeads (Miltenyi). The obtained CD45<sup>neg</sup>CD31<sup>neg</sup> population was then enriched for c-kit<sup>pos</sup> cardiac cells through incubation with anti-human CD117 immunobeads (Miltenyi) (1:10) and then positively sorted using MACS according to the manufacturer's instructions.

Cell Culture

Freshly isolated human c-kit<sup>pos</sup>CD45<sup>neg</sup>CD31<sup>neg</sup> CSCs were plated in CELL-Start (Life Technologies) coated dishes in complete CSC growth medium consisting of a 1:1 ratio of DMEM-F12-Ham's Gibco, Life Technologies) medium and Neurobasal medium (Gibco, Life Technologies), implemented with Embryonic stem-cell FBS (10%, Gibco, Life Technologies), insulin-transferrin-selenium (1%, Life Technologies), epidermal growth factor (20 ng/ml, Peprotech), basal fibroblast growth factor (10 ng/ml, Peprotech), human leukemia inhibitory factor (10 ng/ml, Miltenyi), 37 mg of l-glutamine, B27 supplement (2%, Life Technologies), N2 supplement (1%, Life Technologies), penicillin-streptomycin (1%, Life Technologies), Fungizone (0.1%, Life Technologies) and gentamicin (0.1%, Life Technologies). Cells were maintained in a humidified hypoxic incubator at 37°C, 5% CO<sub>2</sub>, 2% O<sub>2</sub>. Media were replenished every 48 hours and cells were passaged at a 1:4 ratio.
Histology and Immunohistochemistry

Tissue specimens were fixed and embedded in paraffin for histochemical and immunohistochemical analysis. Human tissues were cut in sections of 5 µm. After dewaxing in xylene and rehydration in graded concentration of ethanol, when appropriate, Antigen Retrieval was performed by incubating section in 10 mM sodium citrate buffer (pH 6.0) at 98°C for 30 min. Non-specific antibody binding was blocked by incubation with 10% normal donkey serum (Jackson ImmunoResearch) for 30 minutes at room temperature. Sections were stained for 1h at 37°C or overnight at 4°C and the following primary antibodies were used: c-kit (1:100 dilution, DAKO), c-TnI (1:200 dilution, Abcam), p16INK4A (1:100 dilution, Santa Cruz Biotechnology), 8-OH-dG (1:100 dilution, Origene) and 3-NT (1:400 dilution, Merck Millipore). After washing in phosphate-buffered saline (PBS), sections were incubated with respective secondary antibodies (Jackson ImmunoResearch). Nuclear counterstains were performed by DAPI (4',6-diamidino-2-phenylindole, Sigma) and sections were examined by confocal microscopy (LEICA TCS SP8). The number of positive cells was expressed as a percent fraction of the total cells number per mm².

For immunohistochemical analysis of oxidative stress on tissue sections, after dewaxing, human sections were blocked with Dual Endogenous Enzyme Block for 10 min at room temperature. Then, the sections were stained with antibodies against 8-OH-dG (1:100 dilution, Origene), 3-NT (1:400 dilution, Merck Millipore) and 4-HNE antibody (1:50 dilution, Abcam) for 1h at 37°C. Positive reactions were visualized using a Labelled Polymer-HRP complex and 3,3'-diaminobenzidine tetrahydrochloride (DAB) chromogen using the EnVision+ Dual Link System-HRP (DAKO). Sections were then counterstained with hematoxylin, permanently mounted, and then examined with light microscopy (LEICA, DMI3000B). The morphometric analysis of immunohistochemistry was conducted using Image J.

FACS analysis

Cell analysis was performed on FACSCanto II (BD) with FlowJo software (TREE STAR) to identify the percentage of cardiac small cells expressing different cell-surface markers of interest at passage 2 (P2). A panel of different markers was used for immunophenotypic characterization of human c-kit^pos^CD45^neg^CD31^neg^ CSCs obtained from T2DM and NDM patients. Specific antibodies used are listed in Table 2. Appropriate labelled isotype controls were used to define the specific gates.

Proliferation, clonogenicity, cardiosphere formation and cardiomyocyte differentiation assays in vitro

CSC proliferation was evaluated through BrdU incorporation and growth curve assay, at the indicated time points, on human CSCs. To assess the proliferative activity of freshly isolated c-kit^pos^CD45^neg^CD31^neg^ T2DM-hCSCs and NDM-hCSCs, BrdU 10 μM was administered in vitro. CSCs were plated at density of 1x10⁶ in 24-well CELL-Start-coated dishes and then serum starved in 0% serum base medium. After 48 hrs, starvation medium was replaced by CSC growth medium and BrdU was added to the medium every 6hrs. The cells were fixed at 24 hours and BrdU incorporation was assessed using the
BrdU detection system kit (Roche) according to the manufacturer’s instructions. Nuclei were counterstained with DAPI (Sigma-Aldrich). Cells were evaluated using a fluorescent microscope (LEICA, DMI3000B). Accordingly, the number of BrdU$^{\text{pos}}$ cells was expressed as a percent fraction of the total cell nuclei.

Growth curve assay was archived by plating $5 \times 10^3$ cells in 24-well CELL-Start-coated dishes in CSC growth medium and then serum starved in 0% serum base medium. After 48 hrs, starvation medium was replaced by CSC growth medium and cells were then trypsinized and counted using trypan blue, 1:1 ratio, at the indicated time points.

Single cell cloning was employed through depositing of half c-kit$^{\text{pos}}$CD45$^{\text{neg}}$CD31$^{\text{neg}}$ T2DM-hCSCs and NDM-hCSCs into 96-well CELL-Start-coated Terasaki plates by serial dilution. Individual c-kit$^{\text{pos}}$CD45$^{\text{neg}}$CD31$^{\text{neg}}$ T2DM-hCSCs and NDM-hCSCs were grown in CSC growth medium for 1–3 weeks when clones were identified and expanded. The clonogenicity of the c-kit$^{\text{pos}}$CD45$^{\text{neg}}$CD31$^{\text{neg}}$ T2DM-hCSCs and NDM-hCSCs was determined by counting the number of wells in each 96-well plate containing clones and expressed as a percentage. A total of 10 plates were analyzed.

For cardiosphere generation, $1 \times 10^5$ c-kit$^{\text{pos}}$CD45$^{\text{neg}}$CD31$^{\text{neg}}$ T2DM-hCSCs and NDM-hCSCs were placed in bacteriological dishes with CSC growth medium. Cardiospheres were counted per plate at 14 days and the number expressed as a percentage relative to the number of plated CSCs.

For specific myogenic differentiation, c-kit$^{\text{pos}}$CD45$^{\text{neg}}$CD31$^{\text{neg}}$ T2DM-hCSCs and NDM-hCSCs-derived-cardiospheres were switched to StemPro®-34 SFM differentiation medium (a serum-free medium conditioned with StemPro®-Nutrient Supplement, Gibco, Life Technologies), Glutamine (2mM) and penicillin-streptomycin (1%, Life Technologies). For cardiomyocyte differentiation BMP4 (10ng/ml, Peprotech), Activin-A (50 ng/ml, Peprotech), β-FGF (10ng/ml, Peprotech), Wnt-11 (150ng/ml, R&D System) and Wnt-5a (150ng/ml, R&D System) were added to base differentiation medium. Then differentiating cardiospheres were pelleted and transferred to laminin coated dishes (1µg/ml) and Dkk-1 (150ng/ml, R&D System) was added to base differentiation medium until day 14. Cell differentiation was evaluated at 14 days.

**Immunocytochemistry**

A volume of 200 µl of a $0.15 \times 10^6$ cells/ml suspension of c-kit$^{\text{pos}}$CD45$^{\text{neg}}$CD31$^{\text{neg}}$ CSCs obtained from T2DM and NDM patients were directly loaded in cytofunnel and spin down at 800 rpm for 3 min onto poly-lysine-coated slides using a Shandon Cytospin 4 Cytocentrifuge (Thermo Fisher Scientific). Slides were immediately fixed using PFA 4% (Sigma-Aldrich). After fixation, cells were allowed to air dry before proceeding with immunostaining. Slides were washed with PBS, and incubated in 0.1% Triton X-100:PBS at room temperature for 10 minutes. After washing with 0.1% tween:PBS and 30 minutes in 10% donkey serum, cells were incubated overnight at 4°C with primary antibodies against p16$^{\text{INK4a}}$ (1:100 dilution, Santa Cruz Biotechnology), γ-H2AX (1:400 dilution, Cell Signaling) in 0.1% Tween:PBS. Slides were then
washed and incubated with corresponding secondary antibodies (Jackson Immunoresearch) for 1 hour at 37°C. After washing, the nuclear DNA of the cells was counterstained with DAPI (Sigma-Aldrich) at 1µg/ml and mounted using Vectashield mounting media (Vector labs). Imagines were acquired using a confocal microscope (LEICA TCS SP8). The number of p16^INK4a^ positive and γ-H2AX positive cells was expressed as a percent fraction of the total CSC nuclei. Cell apoptosis were detected using the Terminal deoxynucleotidyl Transferase (TdT) assay (TUNEL assay, Merck Millipore) according to the manufacturer’s instructions. Briefly, to perform TUNEL staining, cells were fixed with paraformaldehyde 2% in PBS (pH 7.4) and permeabilized with 0.1% Triton X-100 in 0.1% sodium citrate. Then cells were incubated with TUNEL reaction mixture (containing Terminal deoxynucleotidyl transferase enzyme + nucleotide mixture in reaction buffer) for 1 hour at 37°C in a humidified atmosphere in the dark. After washing, the nuclear DNA of the cells was counterstained with DAPI (Sigma-Aldrich) and the slides mounted using mounting media (Vector labs). Imagines were acquired using a confocal microscope (LEICA TCS SP8). The number of apoptotic cells was expressed as percentage of the total nuclei.

SA–β-gal staining was performed using a SA–β-gal staining kit (Cell Signaling Technology) according to the manufacturer's instructions and protocols previously reported [23]. In brief, c-kit^pos^CD45^neg^CD31^neg^ T2DM-hCSCs and NDM-hCSCs were grown in culture and then fixed with 1X Fixative Solution for 10-15 min at room temperature. Following fixation, cells were washed with PBS before being incubated in β-Galactosidase Staining Solution (pH 6.0) at 37°C in a dry incubator overnight. The enzymatic reaction was stopped by washing slides with ice-cold PBS and SA–β-gal staining was fixed with ice-cold methanol for 30s before mounting/visualising. Senescent CSCs were identified as blue-stained cells using light microscopy. A minimum of 10 images were taken at x10 magnification from random fields and the percentage of SA–β-gal cells were expressed as percentage of total nuclei.

To perform cTnI staining, human c-kit^pos^CD45^neg^CD31^neg^ CSCs-derived cardiospheres were fixed using PFA 4% (Sigma-Aldrich) after 14 days in differentiation media. After fixation, cells were incubated with cTnI antibody (1:200, Abcam) for 2 hour at 37°C in a humidified atmosphere. The corresponding secondary antibody (Jackson Immunoresearch) was added for 1 hour at 37°C in the dark. The nuclear DNA of the cells was counterstained with DAPI (Sigma-Aldrich) and the slides mounted using mounting media (Vector labs). Imagines were acquired using a confocal microscope (LEICA TCS SP8).

**Telomere length**

Genomic DNA of human c-kit^pos^CD45^neg^CD31^neg^ CSCs obtained from myocardial samples of T2DM and NDM patients was extracted using Quick-DNA Microprep Kit (Zymo Research). Telomere length was analyzed by using the Absolute Human Telomere Length Quantification qPCR Assay Kit (ScienCell Research Laboratories). Genomic DNA (10 ng) was amplified with the FastStart DNA Green Master (Roche Life Science) using a CFX384-Real-Time PCR System (Biorad) and data analysis was conducted according to manufacturer's instruction. For each genomic DNA samples, two different reaction were performed using two primer set: a Telomere primer set to recognize and amplify the telomere sequences and a single copy reference (SCR) primer set to normalize the data that recognizes and amplifies a 100
bp-long region on human chromosome 17, accordingly, a known telomere length as reference to calculate the telomere length of target samples. All reactions were run in triplicate. The average telomere length was calculated by following the manufacturer’s instructions.

Telomerase Activity Quantification

Human c-kit^{pos}CD45^{neg}CD31^{neg} CSCs obtained from myocardial samples of T2DM and NDM patients were processed according to the manufacturer’s protocol using the Telomerase Activity Quantification qPCR Assay Kit (ScienCell Research Laboratories) [24]. Briefly, cell pellets were thawed in lysis reagent enables to release telomerases in the native state. Cell lysate samples were incubated with telomerase reaction buffer at 37°C for 3 hours. qRT-PCR was conducted with FastStart DNA Green Master (Roche Life Science) using a CFX384-Real-Time PCR System (Biorad) and data analysis was performed according to manufacturer’s instruction. The telomere primers set recognizes and amplifies newly synthesized telomere sequences in the assay. All reactions were run in triplicate. Telomerase activity quantification was calculated by following the manufacturer’s instructions.

RNA extraction and RT-PCR analysis

RNA was extracted from human c-kit^{pos}CD45^{neg}CD31^{neg} CSCs obtained from myocardial samples of T2DM and NDM patients, using TRlzol® RNA Isolation Reagents (Ambion) and quantitated using a Nanodrop 2000 Spectrophotometer (Thermo Fisher scientific). Reverse transcription was performed with 0.5-1 µg of RNA using HighCapacity cDNA Kit (Applied Biosystem). Quantitative RT-PCR was performed using TaqMan Primer/Probe sets (see Table 3) using StepOne Plus real Time PCR System (Applied Biosystems). The following genes were tested: GATA-4, NKX2.5, MEF2C TNNT2, ACTC1, MYH6, MYH7, MCP1, IGFBP5, MMP3, IL-6, CCL11, IL-8, GM-CSF and PAI1. Data were processed by the ΔCt method using StepOne Software v2.3 and mRNA was normalized to the housekeeping gene, GAPDH. All reactions were carried out in triplicate.

Western Blot Analysis

Immunoblots were carried out using protein lysates obtained from c-kit^{pos}CD45^{neg}CD31^{neg} T2DM-hCSCs and NDM-hCSCs. Aliquot equivalent of ~40 to 70 µg of proteins were separated on gradient (6–15%) SDS-polyacrylamide gels. After electrophoresis, proteins were transferred onto nitrocellulose filters, blocked with either 5% dry milk and incubated with Ab against p16^{INK4A} (1:1000, Santa Cruz Biotechnology) and GAPDH (1:1000, Santa Cruz Biotechnology). Proteins were detected by chemiluminescence using horseradish peroxidase-conjugated 2Abs and placing the nitrocellulose filters on a photographic film. The acquisition was archived using Medical X-ray processor 2000 (CARESTREAM). Densitometry was obtained using ImageJ software. Immunoblots were performed in biological triplicates.

Cytokine & growth factors assay
Cytokine and growth factor levels [Interleukin-1α (IL-1α), Interleukin-1β (IL-1β), Interleukin-2 (IL-2), Interleukin-4 (IL-4), Interleukin-6 (IL-6), Interleukin-8 (IL-8), Interleukin-10 (IL-10), Interferon γ (IFNγ), tumor necrosis factor alpha (TNFα), Monocyte Chemoattractant Protein-1(MCP-1), Vascular Endothelial Growth Factor (VEGF), Epidermal growth factor (EGF)] were simultaneously evaluated through the “Cytokine & Growth Factors Array (CTK)” kit (Randox Labs) by using Evidence Investigator biochip analyzer (Randox Labs) [25], a chemiluminescent immunoassay in which antibodies bonded on the biochip surface allow the immobilization of a specific analyte, subsequently bonded with a secondary antibody labelled with horseradish peroxidase (HRP), to trigger a luminol-based chemiluminescent signal emission. The light signal generated from each of the test regions is then detected using imaging technology and quantified with a stored calibration curve.

Senolytic drug treatment, viability in vitro

c-kit^{pos}CD45^{neg}CD31^{neg} T2DM-hCSCS were plated in 24-well CELL-Start-coated plates at 40% confluence and left for 72h. Then, the growing medium was replaced with new fresh medium in the absence or presence of a combination of two senolytics drug, Dasatinib (D, LC Laboratories) and Quercetin (Q, Sigma-Aldrich), respectively at doses 0.25μM and 10μM added for 6 hours. Then, the conditioned medium was replaced with complete fresh medium. After 48h the treatment D+Q was repeated as above.

For myogenic differentiation, T2DM-hCSCs were treated or untreated with D+Q in growth media as above. Then cells were placed in bacteriological dishes and the differentiation assay was performed as above described.

Statistical analysis

All data are presented in mean ± standard deviation. Data were analysed using t-test comparisons in GraphPad Prism version 8.0.0 for Windows, GraphPad Software, San Diego, California USA, www.graphpad.com. Differences of p<0.05 are considered statistically significant.

Results

Increased Oxidative Stress in Diabetic Ischemic Cardiomyopathy

Diabetes is characterized by an enhanced oxygen toxicity [26]. ROS is the distal signal of the cascade of events triggered by Diabetes that leads to the initiation of the cell death pathway in the heart [27].

To assess the effects of Diabetes on ROS-mediated cytoplasmic and DNA damage, we obtained left ventricles (LV) samples from patients with Type 2 diabetes mellitus (T2DM) and NON-diabetic mellitus (NDM) patients with post-infarct cardiomyopathy undergoing surgical coronary revascularization. As inclusion criteria an age between 50-64 years was required in order to try separate the effects of aging from Diabetes on senescence. On this premise, we included 10 T2DM and 6 NDM patients from which peri-infarct/border zone biopsies were obtained (Table 1). These samples were analyzed for the presence
of 8-OH-deoxyguanosine (8-OH-dG), Nitrotyrosine (3-NT), and 4-hydroxynonenal (4-HNE). Diabetes status resulted in an increase of myocardial cells, which stained positive for these markers of oxidative stress when compared to NDM patients (Figure 1A-C). In particular, a higher percentage of both cardiomyocytes and CSC-enriched c-kit^{pos}CD45^{neg}CD31^{neg} cardiac cells positive for 8-OH-dG and 3-NT were detected in T2DM as compared with NDM (Figure 2A,B). These data show that cardiac damage in Diabetes is associated with a higher ROS production that targets both muscle cells as well as progenitor cells.

**Increased Oxidative Stress Associated with Increased Expression of Senescent Markers in hCSCs from Diabetic Ischemic Cardiomyopathy**

Cellular senescence is a cell state triggered by stressful insults and certain physiological processes, characterized by a prolonged and generally irreversible cell-cycle arrest with secretory features, macromolecular damage, and altered metabolism. These mechanisms are generally interdependent as indeed oxygen toxicity and DNA damage alter telomeres, resulting in telomere shortening, cellular senescence, and cell dysfunction and/or death [28]. Although a link to organismal aging is clear, aging and senescence are not synonymous as, indeed, cells can undergo senescence, regardless of organismal age [29,30]. On the other hand, aging tissues experience a progressive decline in homeostatic and regenerative capacities, which has been attributed to senescence of their tissue-specific stem cells [31]. On this premise, to assess whether the exaggerated ROS in Diabetes associated with senescence of tissue-resident CSCs, independently of age, we first assessed the expression of p16^{INK4a} in CSCs-enriched c-kit^{pos}CD45^{neg}CD31^{neg} cardiac cells in the myocardial sections from samples obtained from T2DM patients with ischemic cardiomyopathy as compared to NDM patients. p16^{INK4a} inactivates CDK4 and CDK6, which regulate cell-cycle progression in G1; p16^{INK4a} forms binary inhibitor–CDK complexes that are highly stable and cannot be dissociated by cyclins, and cells arrest permanently [32]. Therefore, p16^{INK4a} is widely used as senescent biomarker, despite not being sufficient on its own to define/detect senescent cells [33-36].

The expression of p16^{INK4a} in CSCs-enriched c-kit^{pos}CD45^{neg}CD31^{neg} cardiac cells markedly increased in myocardial samples from T2DM ischemic cardiomyopathy when compared to NDM counterparts (Figure 2C), confirming that a higher oxidative stress couples with the induction of senescence in the cardiac progenitor pool.

To further assess senescence in the adult progenitor pool, we established cultures of human adult cardiac stem cells (hCSCs) from atrial specimens of T2DM (n=6) and NDM (n=6) patients undergoing to cardiac surgery (Table 1) using magnetic activated cell sorting (MACS) for CD45/CD31 negative sorting followed by c-kit positive sorting to isolate CSC-enriched c-kit^{pos}CD45^{neg}CD31^{neg} cardiac cells [12,15,37]. By flow cytometry, the c-kit^{pos}CD45^{neg}CD31^{neg} cardiac cells from T2DM (hereafter T2DM-hCSCs) showed a membrane phenotype typical of cardiac cells enriched with multipotent hCSCs (Figure 3A) as we have previously shown [12], with no significant difference as compared to c-kit^{pos}CD45^{neg}CD31^{neg} cardiac cells from NDM donors (hereafter NDM-hCSCs) (*data not shown*). Indeed, the hCSC-enriched c-
kit^{pos}CD45^{neg}CD31^{neg} cardiac cells at 14 days in culture after their primary harvest, expressed PDGF-Rα, CD166, CD105 and CD90, confirming the enrichment for the phenotypical identity of *bona fide* CSCs (Figure 3A) [37,38].

As first assessment of replicative senescence in the isolate cells, we measured telomerase activity and telomeric length (Figure 3B,C). T2DM-hCSCs showed a significant reduced telomerase activity which associated with a decreased average telomere length when compared to NDM-hCSCs (Figure 3B,C). These data suggest that Diabetes determines exaggerated oxidative stress in hCSCs that is linked to replicative senescence as demonstrated by the appearance of p16^{INK4a} positive cells and reduced telomerase activity inducing telomere shortening.

**CSCs from non-aged diabetic subjects show impaired cell growth and myogenic differentiation potential in vitro**

One common feature of senescent cells is an essentially irreversible cell-cycle arrest that can be an alarm response instigated by deleterious stimuli or aberrant proliferation [30]. The first molecular feature associated with senescence is telomere shortening, a result of the DNA end-replication problem, during serial passages [28,39]. On the basis of the immunohistochemistry data showing increased p16^{INK4a} expression in myocardial sections and of the *in vitro* data demonstrating telomerase deficit with telomere length attrition, we evaluate replicative competence of T2DM-hCSCs. To this aim, freshly-isolated T2DM-hCSCs and NDM-hCSCs were plated for a week and then compared for their growth potential *in vitro*. T2DM-hCSCs show a significant decreased proliferation *in vitro* when compared to NDM-hCSCs, either assessed by growth curve kinetics over time as well as BrdU incorporation over 24 hours (Figure 4A,B). To assess whether the proliferation defect in the expansion capacity of T2DM-hCSCs was coupled with a deficit in other 'stemness' capabilities, we tested these cells for their clonal amplification by depositing half-cell per well in a 96 wells plate and for their spheroid potential by plating them in bacteriological dishes [12]. Remarkably, at 14 days after cell deposition/plating in clonogenic or spherogenic medium, respectively, clonal efficiency was 2.5-fold lower and spherogenesis 2.5-fold lower in T2DM-hCSCs when compared to NDM-hCSCs (Figure 4C,D).

Furthermore, cultured T2DM-hCSCs, expressed biomarkers of cellular senescence at significantly higher levels than NDM-hCSCs. Indeed, T2DM-hCSCs had 10-times higher levels of p16^{INK4a} when compared to NDM-hCSCs (Figure 4E). Cytospin and immunocytochemistry at single cell level show that the number of p16^{INK4a} positive cells within c-kit^{pos}CD45^{neg}CD31^{neg} T2DM-hCSCs (Figure 4F) was significantly higher when compared to NDM-hCSCs (Figure 4G). Concurrently, T2DM-hCSCs show increased senescence-associated beta-galactosidase (b-Gal) positive cells *in vitro* (Figure 4H). Furthermore, γ-H2AX expression revealed that T2DM-hCSCs accumulate significantly more DNA damage than NDM-hCSCs *in vitro* (Figure 4I). Finally, T2DM-hCSCs had a significantly higher apoptotic rate when compared to NDM-hCSCs (Figure 4J).
The above data show that Diabetes increases senescence of hCSCs affecting their expansion, clonal and spheroid potential; we then tested the differentiation potential of these cells as it has been shown in different contexts that senescence and Diabetes both affect differentiation of tissue-dependent stem/progenitor cells [19-21]. Considering that myogenic differentiation is key to myocardial regeneration from endogenous progenitor cells, we evaluated cardiomyocyte differentiation of T2DM-hCSCs vs. NDM-hCSCs when grown in myogenic differentiation medium using the sphere-based myogenic assay that we previously established [38]. RT-PCR data show that T2DM-hCSCs upregulated significantly less than NDM-hCSCs the main cardiac transcription factors (GATA-4, NKX2.5 and MEF2C) and myocyte contractile genes (TNNT2, ACTC1, MYH6 and MYH7) during myogenic differentiation induction (Figure 4K). Only 24±4% of T2DM-hCSCs, as compared to 52±11% of NDM-hCSCs, acquired the prototypical myocyte contractile marker cTnI by 4 weeks in culture, overall revealing a decreased capacity for myogenic differentiation (Figure 4L).

Overall, these data show that Diabetes Mellitus hampers human CSC biology, inducing a variety of hallmarks of senescence that likely contribute to the deficit of their regenerative potential.

**T2DM-hCSCs exhibit a defined senescence-associated secretory phenotype (SASP)**

Senescent cells secrete a plethora of factors, including pro-inflammatory cytokines and chemokines, growth modulators, angiogenic factors, and matrix metalloproteinases (MMPs), collectively known as the senescent associated secretory phenotype (SASP) [8,39]. The SASP constitutes a hallmark of senescent cells and mediates many of their patho-physiological effects, such as reinforcing and spreading senescence in autocrine and paracrine fashions [40], activating immune responses [41], hampering tissue plasticity [42] and contributing to persistent chronic inflammation (known as inflammaging) [43]. Thus, the SASP explain some of the deleterious, pro-aging effects of senescent cells. On the other hand, all these patho-physiological mechanisms are equally active in Diabetes. Therefore, we evaluated whether increased senescence markers and functional defect associated also with SASP in T2DM-hCSCs.

To this aim, we obtained culture media from T2DM-hCSCs and NDM-hCSCs after 24 hours in serum-free conditions to evaluate secretion of several SASP factors, including MMP-3, PAI1, IL-6, IL-8, IL-1β and GM-CSF (Figure 5A,B). Interestingly, T2DM-hCSCs secreted respectively 10-, 7-, 18-, 54- and 56-fold higher amounts of IL-1α, IL-1β, IL-6, MCP-1 and IL-8, when compared to NDM-hCSCs (Figure 5A). Concurrently, mRNA levels of MCP1, IGFBP5, MMP3, IL-6, CCL-11, IL-8, GM-CSF and PAI1 SASP factors were all increased in T2DM-hCSCs vs. NDM-hCSCs (Figure 5B).

We then treated for 7 days NDM-hCSCs with conditioned medium (CM) derived from T2DM-hCSCs (T2DM-CM) and measured cell proliferation and senescence of the treated NDM-hCSCs. As controls, NDM-hCSCs were treated either with normal growth media or CM from parallel cultures of NDM-hCSCs. Treatment of NDM-hCSCs with T2DM-CM resulted in a decreased proliferation (p < 0.05) (Figure 5C,D) and an increased proportion of senescent p16INK4a positive (p < 0.05), SA-β-gal-positive and γ-H2AX-
positive hCSCs when compared to NDM-hCSCs in unconditioned medium (UM) or CM from NDM-hCSCs (Figure 5E).

These data show for the first time that Diabetes induces SASP in hCSCs isolated from non-aged patients. SASP appears to negatively affect neighbor hCSCs whereby senescence begets senescence, suggesting that targeting senescent cells could improve CSC dysfunction in Diabetes.

**Senolytics rescue the regenerative deficit of diabetic hCSCs**

The above data show that T2DM-hCSCs harbor a high fraction of senescent cells that produce and secrete a detrimental SASP. Because senescent cells contribute to the outcome of a variety of cardiac disease, including age-related and -unrelated cardiac diseases like anthracycline cardiotoxicity [5,44-46], much effort has been recently made to therapeutically target detrimental effects of cellular senescence. In regard with the latter, senolytic drugs are agents that selectively induce apoptosis of senescent cells by overriding anti-apoptotic pathways in senescent cells [5]. One of the most well studied senolytics therapeutic approach is the combination of Dasatinib (D) and Quercetin (Q) [5], which we show can target senescent CSCs and improve cardiac function in cardiac aging in mice [5].

To that end, T2DM-hCSCs were plated in 24-well dishes at 40% confluence and left for 2 days. Then, cells were administered with a combination of D+Q at a dose of 0.25 μM D with 10 μM Q. Six hours later, D+Q conditioned medium was replaced with complete fresh medium. Untreated cells served as controls. Two days later, D+Q treatment was repeated as above. Two days after, T2DM-hCSCs were then analyzed for proliferation and markers of senescence, p16\textsuperscript{INK4A}, SA-β-gal and γ-H2AX. As shown in Figure 6, D+Q treatment removed from the cell culture the typical enlarged and flattened cell morphology of senescent cells, consistent with the senolytics combination targeting and removing senescent cells (Figure 6A). Concurrently, D+Q increased T2DM-hCSCs proliferation as evaluated by growth curve kinetics when compared to untreated control T2DM-hCSCs (Figure 6B). Accordingly, D+Q significantly reduced the number of SA-β-gal and γ-H2AX positive T2DM-hCSCs as well as the number of p16\textsuperscript{INK4A} senescent cells, which almost disappeared when compared to untreated T2DM-hCSCs (Figure 6C-E). Furthermore, we determined whether D+Q would abrogate the SASP in T2DM-hCSCs. We found that the level of SASP factors secreted by T2DM-hCSCs was significantly reduced by the administration of D+Q combination (Figure 6F).

Finally, we checked whether D+Q pre-treatment was able to rescue the altered myogenesis potential of diabetic hCSCs. To that end, T2DM-hCSCs were treated or untreated D+Q as above in growth media and then cells were plated in bacteriological dishes for sphere formation [12]. CSC-derived cardiospheres were then plated in laminin for additional 7 days in the stage specific cardiomyogenic media [12]. Interestingly, we found that D+Q pre-treatment restored the myogenic capacity of T2DM-hCSCs as indeed this senolytic combination significantly increased myogenic transcription factor and myogenic contractile gene
expression as compared to untreated cells *in vitro* (Figure 6G). Accordingly, D+Q increased the number of T2DM-hCSC-derived cTnI<sup>pos</sup> cardiomyocytes after 14 days in differentiation media (Figure 6H).

Overall, these findings document that clearance of senescent cells using a combination of D+Q senolytics abrogates the SASP and restores a fully proliferative- and differentiation- competent hCSCs pool in T2DM.

**Discussion**

The main findings emanating from this study are that: i) Ischemic damaged myocardial tissue of T2DM patients is characterized by an exaggerated oxidative stress targeting both cardiomyocytes and cardiac stem/progenitor cells (CSCs); ii) Increased oxidative stress in the myocardium of non-aged T2DM patients associates with an increased number of senescent and dysfucntional T2DM-hCSCs as shown by increased p16<sup>INK4a</sup> expression, reduced telomerase activity and telomere length, reduced proliferation, clonogenesis/spherogenesis and myogenic differentiation; iii) T2DM-hCSCs from non-aged subjects show a senescence-associated secretory phenotype (SASP), as demonstrated by the increased secretion of several SASP factors, including MMP-3, PAI1, IL-6, IL-8, IL-1β and GM-CSF; iv) a combination of two senolytics, Dasatinib and Quercetin, clear senescent T2DM-hCSCs restoring expansion and myogenic differentiation capacities of the diabetic hCSC pool.

T2DM is a pre-conditioning and powerful driver of organismal aging [47]. The biological foundations of aging primarily involve cellular senescence and T2DM is plethoric in senescence-driving factors [47]. An intense debate has existed so far addressing whether senescence precedes or follows the onset of perpetual inflammation and insulin resistance (IR) in T2DM [48]. Independently from “who-precedes-who,” diabetic patients experience an obvious accelerated aging process that increases their susceptibility to morbidity and earlier mortality [49]. Hence, diabetes-affected patients have a significantly shorter life expectancy than non-diabetic individuals, and cardiovascular disease accounts for a large part of the excess mortality [50]. Indeed, adults with T2DM have 2–4 times increased cardiovascular risk compared with adults without diabetes, and the risk rises with worsening glycemic control (T2DM) [1].

Adult tissue-specific stem cells are multipotent cells that are considered a lifelong cellular reservoir to ensure the continuous generation, replacement, and restitution of multiple tissue lineages [51, 52]. Therefore, adult stem cells play a vital role in preventing the aging of organs and tissues, and can delay aging. However, during aging, these cells also undergo some detrimental changes such as alterations in the microenvironment, a decline in the regenerative capacity, and loss of function so that aging has been linked to tissue-specific adult stem cell exhaustion [31]. Converging evidence conclusively demonstrated how toxic high glucose load may be for survival, differentiation plasticity, and regenerative competence for different stem cells lineages [47, 53–57]. The self-sustaining vicious circle amalgamated by hyperglycemia/mitochondrial dysfunction/oxidative stress appears as a master driver to adult mesenchymal stem cell senescence [58]. Accordingly, the diabetic pro-oxidative environment is a major contributing factor for premature adult stem cell senescence and functional deficit [47, 58–62]. Therefore,
Aging and T2DM drive adult stem cell senescence and regenerative deficit and conversely adult stem cell senescence and regenerative deficit drive the progressive pathology in aging and diabetes.

Like other tissue specific adult stem/progenitor cells, also CSCs are not immortal [16]. They undergo cellular senescence characterized by increased ROS production and oxidative stress and loss of telomere/telomerase integrity in response to a variety of physiological and pathological demands with aging [16]. Nevertheless, the old myocardium preserves an endogenous functionally competent CSC cohort which appears to be resistant to the senescent phenotype occurring with aging [16]. The latter envisions the phenomenon of CSC ageing as a result of a stochastic and therefore reversible cell autonomous process, whereby targeting the senescent cells would benefit the recovery of a healthy CSC cohort [16]. Concurrently, T2DM impairs the in vitro proliferative and differentiation potential of human cardiac stem/progenitor cells, worsening their senescence phenotype when compared with CSCs from non-diabetic ischemic patients [22]. Additionally, miR-34a is significantly upregulated while SIRT1 is downregulated in adult CSCs harvested from T2DM patients, which is associated with a higher pro-apoptotic caspase-3/7 activity [63]. All the above studies were, however, conducted in hCSCs isolated from diabetic old patients (> 65 years), which could not clearly distinguish the role of age from DM on hCSC function. On the other hand, in an animal model of insulin-dependent DM in young mice, the myocardial accumulation of ROS drives CSC senescence through the expression of p53 and p16INK4a proteins and telomere erosion [27]. p66shc knockout inhibits CSC senescence and death, preventing the senescent phenotype and the development of cardiac failure by T2DM [27]. Additionally, Diabetes persistently decreases the ability of isolated CSCs from young male mice to proliferate, survive oxidative insults, and differentiate, which can be explained at least in part by an uncoupling of biosynthetic glucose metabolism pathways [64].

Diabetes suppresses CSC activation through a diminished SCF expression in the heart following permanent LAD ligation in young/adult rats [20]. Altogether, these data postulate the tantalizing hypothesis that the premature cellular senescence and ageing of resident CSCs underpins the development of diabetic heart disease [10].

In the present study we show premature aging in hCSCs isolated from non-aged (< 65 years old) patients with T2DM. Indeed, T2DM hCSCs harbor a significant fraction of senescent cells, with exaggerated ROS, DNA damage and telomere erosion. In particular, we provide evidence that independently from age, T2DM-hCSCs have a defined pathologic SASP, which can drive senescence into healthy hCSCs to senescence. Furthermore, T2DM-hCSCs have a reduced capacity to differentiate into cardiomyocytes in vitro. The premature senescence is clearly central to the regenerative defect imposed by Diabetes on hCSCs as demonstrated by the elimination of senescent cells by senolytic treatment, a combination of Dasatinib and Quercetin, which rescued the expansion and myogenic capacity of T2DM-hCSCs. Therefore, this data suggests that eliminating senescent hCSCs by senolytic therapy in vivo could be a therapeutic strategy to prevent hCSC dysfunction even at early stages of Diabetic disease.
Although hampered by the difficulty and limitation of working with samples of human tissue, which makes the data presented here mainly descriptive and falling short of being able to establish direct cause-effect relationship(s) (as it is also the case for most work based on primary human tissue), our findings strongly point to Diabetes hampering human CSC biology, inducing a variety of hallmarks of senescence in non-aged subjects that contribute to the deficit of their regenerative potential. Diabetes induces a pathologic SASP in hCSCs that spread to otherwise healthy hCSCs, reducing the functional pool of hCSCs. This conclusion is reinforced by the evidence that clearance of senescent cells by senolytics abrogates the SASP and restores a fully proliferative- and differentiation- competent hCSC pool in T2DM. Nevertheless, this study cannot fully extrapolate the role of ischemic damage to the observed effects of Diabetes on hCSCs in vivo. Indeed, despite included patients were homogenous for number of vessel diseases and infarct area, Diabetes is known to determine more intense microvascular ischemia, which may have affected the status and function of hCSCs. On the other hand, isolated hCSCs were obtained from right atrial tissue which were not ischemic and therefore most likely the observed effects of Diabetes is independent form ischemia at least for the in vitro data. Yet, further work is needed to fully clarify this issue. Finally, the data of this study in human cells in vitro surely improve our understanding of the cellular mechanisms behind CSC dysfunction in T2DM, identifying senescent cells as potential drug targets and guiding the design of clinical test of candidate D + Q for the myocardial regenerative defect of the diabetic heart. However, human cells in vitro cannot fully recapitulate the human scenario whereby human organoids, stem cell-derived 3D culture systems, re-create the architecture and physiology of human organs in remarkable detail, providing unique opportunities for the study of human disease [65]. It will be therefore truly interesting to further evaluate the role of premature senescence of hCSCs in the context of diabetic human cardiac organoids in the next future [66].

Conclusions

DM hampers human CSC biology, inducing cellular senescence and senescence associate secretory phenotype (SASP) in non-aged subjects (< 65 years) that contribute to a significant deficit of their regenerative potential and in particular to their ability to differentiate in new cardiomyocytes. Clearance of senescent cells by senolytics abrogates the SASP and restores a fully proliferative- and differentiation-competent hCSC pool in T2DM. Therefore, senolytics may represent a potential therapeutic approach to prevent or treat the reduced regenerative potential of the diabetic heart.

Declarations

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Consent for publication: Not Applicable.

Availability of data and materials: All data generated or analysed during this study are included in this published article
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Authors' contributions: Conceptualization: D.T., K.U. and E.C.; Design of the work: A.DeA., G.M.E-H., K.U., D.T., and E.C. Data Acquisition and Analysis: F.M., M.S., L.S., D.C., C.M., N.S. M.G.; Interpretation: D.F., M.T., F.C.S, P.M., A.DeA., K.U., D.T., and E.C. Writing original draft preparation and editing F.M., G.M.E-H., M.R., E.C., K.U., D.T.. All authors have read and agreed to the published version of the manuscript.

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Tables

Table 1. Characteristics of the Patients Enrolled in the Study
Quantitative data are expressed as mean±SD. Binary data are reported by counts. *P<0.01
vs DM patients. Comparisons of the quantitative data have been made through use of
Student's t test for independent samples. The χ² test was used to compare binary data.
T2DM indicates Type 2 Diabetes Mellitus; NDM non diabetes Mellitus; BMI body mass
index; FBG, fasting blood glucose; LDL, low-density lipoprotein; HDL, high-density
lipoprotein; TG, triglycerides; SBP, systolic blood pressure; DBP, diastolic blood pressure;
CHD, coronary heart disease; ACEI, angiotensin-converting enzyme-inhibitor; ARB, angiotensin II receptor blocker.

Table 2. Flow cytometry antibodies
| Antigen          | Antibody ID | Company       | Application |
|------------------|-------------|---------------|-------------|
| c-kit            | A3C6E2      | Miltenyi Biotec | FC          |
| CD184 (CXCR4)   | 12G5        | Miltenyi Biotec | FC          |
| CD9              | SN4 C3-3A2  | Miltenyi Biotec | FC          |
| CD44             | DB105       | Miltenyi Biotec | FC          |
| CD81             | 5A6         | BioLegend     | FC          |
| CD90             | DG3         | Miltenyi Biotec | FC          |
| CD166            | 3A6         | BioLegend     | FC          |
| CD105            | 43A4E1      | Miltenyi Biotec | FC          |
| MDR-1            | UIC2        | BioLegend     | FC          |
| PDGFR-α          | 16A1        | BioLegend     | FC          |
| CD150            | REA151      | Miltenyi Biotec | FC          |
| CD31             | AC128       | Miltenyi Biotec | FC          |
| CD45             | REA747      | Miltenyi Biotec | FC          |
| CD34             | AC136       | Miltenyi Biotec | FC          |
| Mouse IgG1 - Isotype control antibody |             | Miltenyi Biotec | FC          |
| Mouse IgG2a - Isotype control antibody |             | Miltenyi Biotec | FC          |
| Mouse IgG2b - Isotype control antibody |             | Miltenyi Biotec | FC          |

*FC denotes Flow Cytometry*

### Table 3. TaqMan Primers List

| Gene  | Accession  |
|-------|------------|
| GAPDH | Hs02786624_g1 |
| MEF2C | Hs00231149_m1 |
| NKX2.5| Hs00231763_m1 |
| GATA4 | Hs00171403_m1 |
| MYH6  | Hs01101425_m1 |
| MYH7  | Hs01110632_m1 |
| TNNT2 | Hs00943911_m1 |
| ACTC1 | Hs01109515_m1 |
| CCL2  | Hs00234140_m1 |
| CXCL8 | Hs00174103_m1 |
| GM-CSF| Hs00929873_m1 |
| IGF-BP| Hs00181213_m1 |
| IL-6  | Hs00174131_m1 |
| PAI-1 | Hs00167155_m1 |
| MMP-3 | Hs00968305_m1 |
| CCL-11| Hs00237013_m1 |

### Figures
Cardiac damage in diabetes is associated with higher ROS production. (a-c) Light microscopy images show representative DAB-staining of LV samples obtained from T2DM and NDM patients. The respective bar graph show (a) the percentage of 8-OH-dG positive nuclei (in brown), (b) the percentage of 3-NT positive cells (in brown) and (c) the 4-HNE intensity levels (in brown) in T2DM cross section compared to NDM tissue sections (Representative of n=3 heart sections). Scale bar= 200µm. All data are mean±S.D.
Figure 2

Quantification of oxidative stress and senescent markers in myocytes and progenitor cells. (a) Representative confocal images and bar graph showing 8-OH-dG positive cardiomyocytes and 8-OH-dG positive hCSCs in LV samples obtained from T2DM patients (8-OH-dG, red; c-kit, green; cTnI, white; DAPI, blue; representative of n=3 heart sections). Scale bar=50µm and 10µm respectively. (b) Representative light microscopy, bar graph and confocal images showing 3-NT positive cardiomyocytes (3-NT, brown).
and 3-NT positive hCSCs in LV samples obtained from T2DM patients (3-NT, red; c-kit, green; DAPI, blue; representative of n=3 heart sections). Scale bar=100µm and 10µm respectively. (c) Bar graph and representative confocal images showing p16INK4a expression in c-kitposCD45negCD31neg hCSC-enriched cardiac cells in LV samples obtained from T2DM patients compared to NDM counterparts (p16INK4a, red; c-kit, green; DAPI, blue; representative of n=3 heart sections). Scale bar=10µm. All data are mean±S.D.

**Figure 3**

Phenotypic characterization of c-kitposCD45negCD31neg T2DM-hCSCs in vitro. (a) Flow cytometry dot plots showing the membrane phenotype of c-kitposCD45negCD31neg T2DM-hCSCs (Representative of n=3 number of biological replicates) (b,c) Bar graphs showing a telomerase activity (b) and (c) average
Impaired cell growth and myogenic differentiation potential of T2DM-hCSCs in vitro. (a) Cell growth curve shows a decreased proliferation in vitro of c-kitposCD45negCD31neg T2DM-hCSCs compared to NDM-hCSCs (Representative of n=3 number of biological replicates). (b-d) Bar graphs showing respectively
BrdU incorporation, clonogenesis and spherogenesis in c-kitposCD45negCD31neg T2DM-hCSCs compared to NDM-hCSCs (Representative of n=3 number of biological replicates). (e) Representative Western Blot showing p16INK4a levels in T2DM-hCSCs compared to NDM-hCSCs (Representative of n=3 biological replicates). (f) Flow cytometry dot plots showing a hCSCs population obtained from cardiomyocyte-depleted cardiac cell preparations. Left, representative gating strategy using isotype antibodies. Right, post-sorting flow cytometry dot plots showing c-kit expression in the cardiomyocyte-depleted cardiac cell preparations negatively sorted for CD45 and CD31 (Representative of n=3 experiments). (g) Bar graph and representative confocal microscopy images from cytospin preparations showing p16INK4a expression (red) in c-kitposCD45negCD31neg T2DM-hCSCs compared to NDM-hCSCs. Bar graph=75µm (Representative of n=3 biological replicates). (h) Bar graph and representative light microscopy images showing senescence-associated β-Gal positive cells (blue) in T2DM-hCSCs compared to NDM-hCSCs. Scale bar=200µm. (i,j) Bar graphs and representative confocal microscopy images from cytospin preparations of c-kitposCD45negCD31neg T2DM-hCSCs and NDM-hCSCs showing (i) the expression of γ-H2AX, (green) and (j) TdT (green) in T2DM-hCSCs compared to NDM-hCSCs. Bar graph=75µm and 50µm. (Representative of n=3 biological replicates). (k) Heat maps showing qRT-PCR analysis of the main cardiac transcription factors and myocyte contractile genes in c-kitposCD45negCD31neg T2DM-hCSCs compared with NDM-hCSCs 14 days after myogenic differentiation induction. Color scale indicates change in Ct (threshold cycle) relative to the normalized GAPDH control (Representative of n=3 biological replicates). (l) Bar graph showing cTnI expression levels of differentiated T2DM-hCSCs compared to NDM-hCSCs (Representative of n=3 biological replicates). All data are mean±S.D
The senescence-associated secretory phenotype (SASP) of T2DM-hCSCs. (a) Bar graphs showing the SASP factor protein levels in culture media from T2DM-hCSCs and NDM-hCSCs after 24 hours in serum-free conditions (Representative of n=3 biological replicates). (b) Bar graphs showing transcript SASP factors expression in T2DM-hCSCs vs. NDM-hCSCs (Representative of n=3 biological replicates). (c) Cell growth curve showing the in vitro proliferative change of NDM-hCSCs placed in conditioned medium (CM) derived from T2DM-hCSCs compared to the NDM-hCSCs placed in unconditioned medium (UM) and NDM-hCSCs CM (Representative of n=3 number of biological replicates). (d) Bar graph showing BrdU incorporation of NDM-hCSCs placed in UM, NDM-CM and T2DM-CM (Representative of n=3 biological replicates). (e) Bar graphs showing senescent p16INK4a positive hCSCs, SA-β-gal-positive hCSCs and γ-H2AX-positive hCSCs in NDM-hCSCs treated with T2DM-CM compared to UM and NDM-CM (Representative of n=3 number of biological replicates). All data are mean±S.D.
Senolytics ameliorates regenerative deficit of diabetic hCSCs. (a) Light microscopy images showing that the typical enlarged and flattened morphology of senescent cells present in untreated control (CTRL) disappear in D+Q treated T2DM-hCSCs in vitro. Scale bar=400μm (b) Cell growth curve shows the in vitro proliferation of c-kitposCD45negCD31neg T2DM-hCSCs after D+Q treatment compared to CTRL per se (Representative of n=3 number of biological replicates). (c-e) Bar graphs showing the number of (c) SA-β-gal, (d) p16INK4a senescent cells and (e) γ-H2AX positive cells in T2DM-hCSCs after D+Q treatment compared to CTRL (Representative of n = 3 number of biological replicates). (f) Bar graphs showing transcript SASP factors expression in T2DM-hCSCs after D+Q treatment vs. CTRL (Representative of n=3 biological replicates). (g) Heat maps showing qRT-PCR analysis of the main cardiac transcription factors and myocyte contractile genes (GATA-4, NKX2.5, MEF2C, TNNT2, ACTC1, MYH6 and MYH7) in
differentiating T2DM-hCSCs after D+Q treatment vs. CTRL. Color scale indicates change in Ct (threshold cycle) relative to the normalized GAPDH control (Representative of n=3 biological replicates). (h) Bar graph show cTnl expression levels in differentiated T2DM-hCSCs after D+Q treatment vs. CTRL (Representative of n=3 biological replicates). All data are mean±S.D