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To cite this version:
Marion de Toledo, Isabel Lopez-Mejia, Patricia Cavelier, Marine Pratlong, Célia Barrachina, et al.. Lamin C Counteracts Glucose Intolerance in Aging, Obesity, and Diabetes Through β-Cell Adaptation. Diabetes, American Diabetes Association, 2020, 69 (4), pp.647-660. 10.2337/db19-0377. hal-03065207

HAL Id: hal-03065207
https://hal.archives-ouvertes.fr/hal-03065207
Submitted on 14 Dec 2020

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Lamin C counteracts glucose intolerance in aging, obesity and diabetes through β-cell adaptation

Running title: Lamin C involvement in β-cell adaptation

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Total word count : 5293 words
Number of figure/table: 6 figures, 6 supplemental datas, 2 supplemental tables
Abstract

Aging-dependent changes in tissue function are associated with the development of metabolic diseases. However, the molecular connections linking aging, obesity and diabetes remain unclear. Lamin A, lamin C and progerin, products of the Lmna gene, have antagonistic functions on energy metabolism and lifespan. Lamin C, albeit promoting obesity, increases lifespan suggesting that this isoform is crucial for maintaining healthy conditions under metabolic stresses. Since β-cell loss during obesity or aging leads to diabetes, we investigated the contribution of lamin C to β-cell function in physiopathological conditions. We demonstrate that aged lamin C-only expressing mice (Lmna^{LCS/LCS}) become obese but remain glucose tolerant, due to adaptive mechanisms including increased β-cell mass and insulin secretion. Triggering diabetes in young mice revealed that Lmna^{LCS/LCS} animals normalize their fasting glycemia by both increasing insulin secretion and regenerating β-cells. Genome-wide analyses combined to functional analyses revealed an increase of mitochondrial biogenesis and global translational rate in Lmna^{LCS/LCS} islets, two major processes involved in insulin secretion. Altogether, our results demonstrate for the first time, that the sole expression of lamin C protects from glucose intolerance through a β-cell adaptive transcriptional program during metabolic stresses highlighting Lmna gene processing as a new therapeutic target for diabetes treatment.
Introduction

Lamin A, lamin C and progerin proteins, produced by alternative RNA processing of the \textit{Lmna} gene, are intermediate filaments that bridge peripheral heterochromatin to the nuclear envelope (1), thus participating in the maintenance of chromatin architecture (2,3) and in transcriptional regulation (4). Mutations in the \textit{Lmna} gene result in laminopathies, including progeria, which lead to metabolic abnormalities such as lipodystrophy, hypertriglyceridemia, insulin resistance and metabolic syndrome, thus suggesting an important role for A-type lamins in the regulation of glucose and lipid metabolism (5). Comparing Knock-in mice expressing all three lamin isoforms (lamin A, lamin C and progerin) or exclusively lamin C isoform (\textit{Lmna^{LCS/LCS}}), we previously reported antagonistic functions of \textit{Lmna} isoforms in energy expenditure and lifespan, pointing out a key function for this gene in the control of energy homeostasis (6). Despite showing increased body weight, \textit{Lmna^{LCS/LCS}} have an unexpectedly increased lifespan suggesting a role for lamin C in improving metabolic health during aging.

Pancreatic β cells play a critical role in maintaining glucose homeostasis through the secretion of insulin, and their dysfunction results in diabetes. Whereas type 1 diabetes (T1D) is an autoimmune disease that results from the destruction of β-cells, type 2 diabetes (T2D) is due to energy imbalance and is characterized by impaired peripheral insulin sensitivity followed by defects in insulin secretion. The common hallmark of T1D and T2D is an inadequate supply of functional insulin-producing β-cells, leading to the dysregulation of glucose metabolism and associated metabolic complications (7). The restoration of normal β-cell mass and function has therefore become of great interest for the generation of novel anti-diabetic drugs. In this context, β-cell regeneration has been extensively explored in mice and humans. However, the mechanisms involved in the functional plasticity required for the regeneration of β-cell mass remain poorly understood (7, 8).
Although recent reports highlighted that \textit{Lmna} isoforms are involved in the structure and function of the exocrine pancreas (9), their contribution to the endocrine pancreas and, more particularly, in insulin-producing $\beta$-cells remain to be investigated. We hypothesized that lamin C might contribute to maintain adequate glucose homeostasis during aging through a role on pancreatic $\beta$-cell plasticity or regeneration. Using a comprehensive analysis of the metabolic phenotype of \textit{Lmna}^{LCS/LCS} mice using different models related to $\beta$-cell failure (\textit{i.e.} aging, STZ-induced diabetes and obesity-induced diabetes), we demonstrate that exclusive lamin C expression favors $\beta$-cell mass, insulin production and insulin secretion in conditions where dual expression of lamin A and lamin C from the \textit{Lmna} gene are accompanied by abnormal glucose homeostasis. This adaptive effect on $\beta$-cell mass and function is associated with an increase in mitochondrial activity and of global translation rates, providing a molecular link between lamin C and $\beta$-cell fitness during aging. Using a model of T2D (diet-induced obesity, DIO), we report that \textit{Lmna}^{LCS/LCS} mice enhance insulin secretion to adapt to DIO and to maintain normoglycemia. Finally, in streptozotocin-induced diabetes mice $\beta$-cell failure was rescued in \textit{Lmna}^{LCS/LCS} through increased regeneration of $\beta$-cells. Altogether, our findings demonstrate that the exclusive expression of lamin C improves $\beta$-cell function and mass during metabolic stress and suggest that the manipulation of \textit{Lmna} isoforms can be targeted to counteract diabetes.
Research Design and Methods

Animal experiments and ethics statement

_Lmna^{LCS/LCS}_ Knock-In mice were bred onto C57BL/6J background as described (10). All animal procedures were conducted in accordance with the French national animal care guidelines. Mice were maintained in pathogen-free conditions in our animal facility (E34-172-16). Mice were housed under a 12-h light/dark cycle and given an *ad libitum* regular chow diet (A04; Safe). All experiments were conducted by authorized personnel (agreements MDT34-433; CC-I-34UnivMontp-f1-12) and approved by the Ethics Committee of the Languedoc-Roussillon region Montpellier (CEEA-LR-12003). Male mice were used for _in vivo_ experiments and isolated islets were prepared from male or female mice.

**Histology and immunohistochemistry**

Pancreas sections (4µm) were deparaffinized, rehydrated, stained with hematoxylin and eosin (HE) and visualized with a Nanozoomer. Islets diameter were measured using NDP View software.

For immunohistochemistry, rehydrated sections were incubated in citrate buffer at 95°C for 30min. After blocking, sections were incubated in primary antibodies (see table 2) overnight at 4°C and were detected using appropriate secondary antibodies. Nuclei were counterstained with Hoechst® 33342.

**Metabolic analysis and generation of diabetic mouse models**

For IPGTT, 18 h-fasted mice were injected intraperitoneally with glucose (1.7g/kg), and glycemia was measured using a blood glucometer (Roche). Insulin levels were quantified using
Mouse Ultrasensitive Insulin ELISAs (ALPCO). To induce T2D, 12-week-old mice had unrestricted access to a HFD (ResearchDiet) for 25 weeks. To induce acute β-cell loss, 13 weeks old mice were injected i.p. with a single high dose (200mg/kg) of streptozotocin (Sigma-Aldrich) dissolved in 0.1M sodium citrate buffer (pH4.4).

**Pancreatic islet Isolation and Glucose-stimulated insulin secretion**

Mouse islets were isolated by type V collagenase digestion (Sigma) of pancreata. After separation in a density gradient medium (Histopaque®-1119, Sigma), islets were handpicked. They were then cultured for 18–20h at 37°C in a 95% air-5% CO$_2$ atmosphere in RPMI-1640 (ThermoFisher) containing 10% FCS and 100 µg/ml penicillin-streptomycin.

To evaluate insulin secretion, 3 replicates of 5 islets were preincubated for 1h at 37°C in KRB buffer. After low (3mM) or high (20mM) glucose stimulation, secreted insulin and total insulin cell content were measured. Data are expressed as a ratio of secreted insulin/total insulin content.

**Total pancreatic insulin and glucagon content**

Pancreata were lysed in 2ml Acid/Ethanol lysis buffer and incubated O/N at 4°C. After neutralization with 10% of 1M Na2CO3, total insulin and glucagon content were measured by ELISA (Mouse Ultrasensitive Insulin elisa-ALPCO, Mercodia Glucagon ELISA - 10 μL).

**RT-qPCR analysis and mitochondrial DNA quantification**

Total RNA was extracted from isolated islets using TRIzol (Life Technologies, USA). Samples were quantified with a NanoDrop ND-1000. 1.5 µg RNA were then converted to cDNA
with the Maxima First Strand cDNA Synthesis Kit (#K1642, Thermo Scientific). Gene expression was normalized to cyclophillin and TBP.

For mitochondrial DNA quantification, islets were digested using proteinaseK. qPCR analysis was performed with primers specific for mitochondrial genes (*mtND5, mt16s*) and normalized to nuclear genes (*LN11, HK2*).

qPCR was performed with a LightCycler® 480 and LightCycler® 480 SYBR GreenI Master (Roche). Primer sequences are provided in Supplemental Table 1.

**Ultrastructural transmission electron microscopy**

After isolation, 150 islets were immersed in a solution of 2.5% glutaraldehyde. After dehydration, samples were embedded in EmBed812. 70nm sections were counterstained with uranyl acetate and observed on a Hitachi-7100 microscope.

**Seahorse experiments and metabolic flux analysis**

Oxygen consumption rate (OCR) was measured in 96-well plates using a XFe96-Seahorse (Agilent). Briefly, 25 islets/well were plated in poly-D-lysine-coated wells in XF-assay medium containing 3mM glucose, 1mM sodium pyruvate and 1mM glutamine (pH7.4) for 1 h at 37°C without CO$_2$. OCRs were monitored in response to glucose (20mM), oligomycin (2μM), FCCP (1μM) and rotenone (0.5μM). OCRs measurements were normalized by total DNA level using a Qubit dsDNA HS assay kit (LifeTechnologies).

**Incorporation of $^{35}$S-Met/Cys into isolated islets**

Seventy isolated islets were incubated in Met/Cys-depleted RPMI medium (Sigma) for 30min and then supplemented with 50μCi $^{35}$S-Met/Cys EasyTag Express (PerkenElmer) for
30min at 37°C. Islets were lysed in RIPA buffer. 10µg of total protein were size-fractionated on 4-12% gradient SDS-PAGE. The gel was dried and quantified using a Phospholmager (GE-Healthcare).

**Flow cytometry**

Islets dissociated with 0.05% Trypsine-0.5 mM EDTA were fixed with 1% PFA and permeabilised with 0.05% Triton X-100. After blocking, cells were incubated anti-insulin antibody, washed, reacted with a FITC-conjugated antibody and analyzed by FACSCANTO II (Becton Dickinson).

**RNA isolation, RNA-seq, RRBS, ATAC-seq and data analysis**

Total RNA was extracted using TRizol and quantified on Qubit (Life Technologies). RNA quality was verified using RNA pico-chips on a 2100-Bioanalyzer (Agilent). Total DNA was purified using MasterPure™ Complete DNA (Illumina). Libraries for RNA-seq and RRBS were prepared and sequenced by the MGX sequencing platform in Montpellier (France).

For ATAC-seq analyses, islets were cryopreserved in 50%FCS, 40%RPMI, 10%DMSO and processed by ActiveMotif.

All data are deposited in GEO-NCBI under accession number GSE122361.

**Statistical analysis**

Results were expressed as the mean ± SEM. Statistical comparison were made using student’s t-test. Statistically significant differences are indicated as *p<0.05, **p< 0.01, ***p< 0.001, ****p< 0.0001
A detailed description of the all procedures and bioinformatic analyses can be found in Supplementary Methods.

**Data and Resource Availability**

The resources generated during and/or analyzed during the current study are available from the corresponding authors on reasonable request. The genome wide dasas generated and/or analyzed during the current study are deposited in GEO-NCBI under accession number GSE122361.
Results

Increased obesity in aged \( Lmna^{\text{LCS/LCS}} \) mice is associated to improved glucose tolerance

To evaluate whether lamin C regulates \( \beta \)-cell function, we evaluated glucose homeostasis in \( Lmna^{\text{LCS/LCS}} \) mice (6). Initially designed to generate a mouse model of progeria, the c.1827C>T;p.G609G point mutation was introduced into a floxed neo cassette (Supplemental data 1A). This cassette was inserted in intron-10 of the \( Lmna \) gene, downstream of the polyadenylation site of lamin C, thus preventing the formation of prelamin A transcripts by blocking lamin A-specific splicing (10). The conditional mutant allele (\( Lmna^{\text{LCS}} \)) resulted in the exclusive expression of lamin C. This was confirmed by comparing its levels in brain, liver, pancreas and muscle from \( Lmna^{+/+} \) (wild type, WT) and \( Lmna^{\text{LCS/LCS}} \) mice (Supplemental data 1B). We observed increased expression of lamin C in \( Lmna^{\text{LCS/LCS}} \) mice compared to WT in most tissues except in the brain, where the lamin C/lamin A ratio is already high even in WT mice, as previously shown (11).

With the experimental \( Lmna^{\text{LCS/LCS}} \) model established, we then investigated the metabolic phenotype of these mice. The \( Lmna^{\text{LCS/LCS}} \) mice gained more weight than control mice as they aged (Fig 1A). This increase in body weight is mainly due to an increase in white adipose tissue without any further change in body composition as shown in our previous study (6). Body weight and fasting glucose levels in \( Lmna^{+/+} \) and \( Lmna^{\text{LCS/LCS}} \) mice were similar until 25 weeks of age (Fig 1A-B). Interestingly, the significant increase in body weight was observed from 30 weeks of age in \( Lmna^{\text{LCS/LCS}} \) mice, but was even more pronounced at 75 weeks of age. The following aging experiments are thus conducted by comparing “young” (25 weeks old) to “old” animals (75 weeks old). We observed a significant increase in fasting glycemia in control mice as they aged (Fig 1B). Interestingly \( Lmna^{\text{LCS/LCS}} \) mice remained normoglycemic, although an increased body weight is typically associated with hyperglycemia. These findings prompted us
to analyze glucose homeostasis in aged \( \text{Lmna}^{\text{LCS/LCS}} \) mice. Glucose clearance was similar in young animals of both genotypes (Fig 1C, left panel), whereas old \( \text{Lmna}^{\text{LCS/LCS}} \) mice showed improved glucose tolerance compared to age-matched \( \text{Lmna}^{+/+} \) mice (Fig 1C, right panel). As expected, no differences in basal and glucose-stimulated insulin secretion were observed in young animals from either genotype (Fig 1D, left panel). Basal and stimulated insulin secretion was, however, increased in old \( \text{Lmna}^{\text{LCS/LCS}} \) mice compared to old WT mice (Fig 1D, right panel). Altogether, these data suggest that although \( \text{Lmna}^{\text{LCS/LCS}} \) mice become obese as they age, their glucose clearance and insulin secretion are improved, suggesting that lamin C may modulate the function of pancreatic islets of Langerhans during aging.

**\( \text{Lmna}^{\text{LCS/LCS}} \) mice have increased β-cell mass and insulin secretion during aging**

Our observations in aged \( \text{Lmna}^{\text{LCS/LCS}} \) mice prompted us to study whether lamin C could contribute to pancreatic islet function by regulating pancreatic mass and/or insulin secretion. Old \( \text{Lmna}^{\text{LCS/LCS}} \) mice had heavier pancreata, whereas no differences in mass were observed between genotypes in young animals (Fig 2A). Histological analyses revealed that while young \( \text{Lmna}^{+/+} \) and \( \text{Lmna}^{\text{LCS/LCS}} \) pancreata were microscopically similar, pancreata from old \( \text{Lmna}^{\text{LCS/LCS}} \) mice displayed significantly larger islets of Langerhans compared to \( \text{Lmna}^{+/+} \) mice (Fig 2B). This effect was further amplified in the pancreata of “very old” (141 weeks old) \( \text{Lmna}^{\text{LCS/LCS}} \) mice, as evidenced by hyperplasic islets (Supplemental data 1C). In young animals, measurements of islet size (Fig 2C, upper panel) and staining of insulin and glucagon-positive cells within the pancreatic islets (Fig 2D, upper panel) revealed no differences, whereas old \( \text{Lmna}^{\text{LCS/LCS}} \) animals displayed enlarged islets with an increase in insulin-positive cells, but no differences in the number of glucagon-positive cells (Fig 2C and 2D, lower panel). Quantification of the \( \beta-/\alpha \)-cell ratio confirmed an increased number of insulin-positive \( \beta \)-cells in aged \( \text{Lmna}^{\text{LCS/LCS}} \) mice (Fig 2E).
whereas no changes were observed on glucagon-positive α-cells (Supplemental data 2A). In addition, when compared to young mice, old \( Lmna^{ LCS/ LCS} \) mice exhibited an increased number of islets per pancreas (Fig 2F). Furthermore, total pancreatic insulin level was increased in LCS mice compared to control mice (Fig 2G) whereas global glucagon level was not significantly different (supplemental data 2B). These data demonstrate that the expression of lamin C protects from aging-induced β-cell mass loss by increasing the size and the number of islets of Langerhans and the number of insulin-positive cells.

We then investigated the functionality of these abundant and large \( Lmna^{ LCS/ LCS} \) islets through glucose-stimulated insulin secretion (GSIS) experiments. Although no difference in GSIS between genotypes was observed in pancreatic islets isolated from young animals, islets derived from old \( Lmna^{ LCS/ LCS} \) mice displayed a 29.5% higher GSIS than age-matched control islets after 20mM glucose stimulation (Fig 2H). Interestingly, no difference of insulin secretion in basal, 2.8mM glucose stimulation was observed, suggesting that the increased insulinemia observed in fasted \( Lmna^{ LCS/ LCS} \) mice (Fig 1D) could be due to a cumulative effect of the increased number of islets and their improved insulin secretion capacity.

To confirm the direct effect of lamin C on β-cells, we next generated MIN6 pancreatic β-cell lines in which lamin A expression was abolished using CRISPR/Cas9-mediated genome editing (MIN6\( ^{ LCO} \)). Targeting exon 11 of the \( Lmna \) gene resulted in a complete depletion of lamin A expression. In the MIN6\( ^{ LCO} \) cells expressing only lamin C (Supplemental data 3A) insulin secretion in response to a 20mM glucose stimulation was increased compared to the control MIN6\( ^{ WT} \) cells (Supplemental data 3B), indicating that lamin C can enhance insulin secretion and improve β-cell function.

Taken together, these results showed that old \( Lmna^{ LCS/ LCS} \) mice have increased β-cell number and improved insulin secretion in response to glucose both \textit{in vitro} and \textit{in vivo}. These
adaptive mechanisms may account for the normoglycemia observed in old $Lmna^{LCS/LCS}$ mice despite their obesity.

**Increased insulin secretion in $Lmna^{LCS/LCS}$ mice can counteract obesity-induced type 2 diabetes**

To counteract obesity, $\beta$-cells need to increase their secretory capacity by progressive expansion of their mass (12). To evaluate how $Lmna^{LCS/LCS}$ $\beta$-cell function adapts to metabolic challenges, we investigated the capacity of $Lmna^{LCS/LCS}$-derived islets to counteract hyperglycemia observed during obesity-induced T2D in young mice. Challenging 12 weeks old $Lmna^{+/+}$ and $Lmna^{LCS/LCS}$ mice with a 60%kcal high-fat diet (HFD) for 25 weeks resulted in increased body weight (Fig 3A). This body weight gain was significantly higher in $Lmna^{LCS/LCS}$ mice and was associated to increased food intake when compared to the control mice (Supplemental data 4). No differences were observed between genotypes in the consumption of regular chow diet (CD). Monitoring of fasting glycemia over the course of HFD revealed that during the first 10 weeks, $Lmna^{LCS/LCS}$ mice were hyperglycemic compared to control mice, probably due to their higher body weight (Fig 3B). However, at week 15 under HFD, a significant improvement in fasting glycemia was observed. At week 25, $Lmna^{LCS/LCS}$ mice demonstrated a further improvement in glucose tolerance (Fig 3C) and showed increased insulin secretion in response to in vivo glucose stimulation (Fig 3D). GSIS experiments using islets isolated from HFD-fed mice for 25 weeks demonstrated that while $Lmna^{+/+}$-derived islets failed to secrete insulin in response to glucose, islets from HFD-fed $Lmna^{LCS/LCS}$ mice efficiently secreted insulin in response to glucose (Fig 3E).

These results demonstrate that $Lmna^{LCS/LCS}$ mice efficiently counteract HFD-induced hyperglycemia by increasing insulin secretion. Taken together with our aging studies, these
results suggest that a lamin C-specific adaptive mechanism is engaged by insulin-producing cells to maintain glucose responsiveness in order to counteract the effects of aging and obesity-induced T2D.

**Increased β-cell regeneration in streptozotocin-injected Lmna\textsuperscript{LCS/LCS} mice**

To further explore the role of lamin C on β-cell number and function, we evaluated the capacity of Lmna\textsuperscript{LCS/LCS} mice to regenerate β-cells in an acute β-cell loss context. 13 weeks old Lmna\textsuperscript{+/+} and Lmna\textsuperscript{LCS/LCS} animals were treated with a single dose of streptozotocin (STZ, 200 mg/kg) to mimic diabetes, and the regeneration of β-cells was measured 18 days after STZ injection. 3 days after STZ treatment, Lmna\textsuperscript{+/+} and Lmna\textsuperscript{LCS/LCS} mice became diabetic with no significant difference in fasting glucose level (Fig 4A). From day 4, while Lmna\textsuperscript{+/+} mice remained hyperglycemic, blood glucose levels in Lmna\textsuperscript{LCS/LCS} mice rapidly decreased with a complete normalization of glycemia 18 days after STZ injection. Moreover, we observed approximately 15% of body weight loss during the first 4 days following STZ treatment for both genotypes. 18 days after STZ injection Lmna\textsuperscript{+/+} mice remained as leaner as they were before the treatment, while Lmna\textsuperscript{LCS/LCS} mice recovered their initial body weight (Supplemental data 5A). Consistent with our previous observations, Lmna\textsuperscript{LCS/LCS} showed improved glucose tolerance 18 days after STZ injection (Fig 4B), suggesting functional insulin-producing cells in STZ-treated Lmna\textsuperscript{LCS/LCS} mice.

Morphological analysis of the pancreata 18 days post-STZ injection revealed that the number of insulin-positive cells was dramatically reduced in Lmna\textsuperscript{+/+} islets (Fig 4C, left panel). Conversely, Lmna\textsuperscript{LCS/LCS} insulin-positive β-cells recovered efficiently, with comparable staining of insulin- and glucagon-positive cells before and after STZ treatment. Overall, 18 days after STZ
injection, a 2.5-fold increase in insulin-positive cells was observed in the pancreas of $Lmna^{LCS/LCS}$ mice compared to control mice (Fig 4C, right panel).

The increased number of islets in $Lmna^{LCS/LCS}$ mice led us to hypothesize that $Lmna^{LCS/LCS}$ mice are either protected from STZ-induced apoptosis and/or are able to efficiently regenerate β-cells after STZ-induced damage. 72h after STZ treatment, β-cells were strongly depleted equally in $Lmna^{+/+}$ and $Lmna^{LCS/LCS}$ mice as shown by representative immunostaining as well as β-cells mass quantification (Fig 4D). The susceptibility of β-cells to STZ-induced apoptosis was confirmed by immunostaining (Supplemental data 5B, left panel), and quantitative assessments 48h after STZ treatment using TUNEL assays showed no significant difference in the number of apoptotic cells between $Lmna^{+/+}$ and $Lmna^{LCS/LCS}$ mice (Supplemental data 5B, right panel). This finding was further confirmed in the MIN6 β-cell line treated with STZ, in which no differences in the number of apoptotic cells were observed between MIN6WT and MIN6LCO cells (Supplemental data 5C).

We next studied β-cell regeneration after STZ treatment in $Lmna^{LCS/LCS}$ mice. The presence of proliferating cells after STZ treatment was quantified using phospho-histone H3 (phosho-H3) antibody staining on pancreatic sections at day 9 post-STZ injection. Pancreata from $Lmna^{LCS/LCS}$ mice had a significantly higher number of phosho-H3 positive cells compared to control mice (Fig 4E), suggesting an increased regenerative capacity in $Lmna^{LCS/LCS}$ mice. To identify the functional profile of these newly generated cells, we next performed co-immunostaining for phosho-H3 (red) and different pancreatic markers, including insulin (β-cells), lectin (ductal cells), glucagon (α-cells) and somatostatin (δ-cells). Co-immunostaining was observed for phosho-H3 and insulin, as well as lectin but not with glucagon or somatostatin, suggesting that proliferating cells observed in $Lmna^{LCS/LCS}$ mice may arise from duplication of
pre-existing β-cells and from ductal cells (Fig 4F) two well-known processes for β-cell neogenesis (8).

Altogether, these results suggest that \( Lmna^{LCS/LCS} \) mice are able to counteract STZ-induced diabetes by regenerating β-cells.

\( Lmna^{LCS/LCS} \) mice display a transcriptional reprogramming of genes involved in mitochondria and translation

Given the important role of lamins in chromatin organization and in the control of gene expression (4), we next evaluated the transcriptional status, DNA methylation and chromatin accessibility of islets isolated from 75 weeks old \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice. RNA-sequencing based analysis (RNA-seq) revealed that 1507 genes (9.3% of the sequenced genes) were significantly differentially regulated between genotypes (p-value < 0.05), with 1024 genes upregulated and 483 genes downregulated in \( Lmna^{LCS/LCS} \) mice relative to controls (Fig 5A). Gene ontology enrichment analysis showed an enrichment of genes associated with mitochondria (15.92%) and global translation (15.59%). Most genes associated with mitochondrial function (205/240 genes) and translation pathways (169/235 genes) were significantly upregulated in \( Lmna^{LCS/LCS} \) islets compared to \( Lmna^{+/+} \) islets. Reduced representation bisulfite sequencing (RRBS)-based methylome analysis revealed that the observed variations in gene expression were not correlated to changes in DNA methylation (Fig 5B). Indeed, independent of the region of the genome that was analyzed (Supplemental data 6), no differentially methylated cytosines (DMC) were found in 1383868 cytosines tested. Moreover, no differentially methylated regions (DMR) were found in 872305 tested regions (GEO accession number: GSE122361). In order to evaluate whether the transcriptional dysregulation was due to a global remodelling of the chromatin compaction and/or accessibility, we performed Assay for Transposable-Accessible
Chromatin with high-throughput sequencing (ATAC-seq) on pancreatic islets isolated from old Lmna^{+/+} vs Lmna^{LCS/LCS} mice. ATAC-seq results revealed that no difference in chromatin accessibility profiles were observed between Lmna^{+/+} and Lmna^{LCS/LCS} isolated islets (Fig 5C), suggesting that the transcriptional dysregulations observed in RNA-seq analysis were not due to changes in global chromatin accessibility upon the exclusive expression of lamin C.

To go further into the molecular mechanisms by which lamins could modulate transcription programs, we analyzed the chromatin architecture by investigating the spatial nuclear localization of selected histone marks that are known to modulate gene expression (13). Immunofluorescence analyses of pancreatic sections showed an enrichment of the active enhancer mark H3K27Ac at the nuclear periphery in Lmna^{LCS/LCS} β-cells compared to control (Fig 5D, left panel) whereas no difference was observed for the localization of the repressive mark H3K9Me3 (Fig 5D, right panel). Interestingly, the distribution of the H3K27Ac epigenetic mark to the nuclear periphery is associated with a modification of the nuclear architecture. Indeed, we observed that the chromatin is relocated to the periphery of the nucleus in Lmna^{LCS/LCS} mice as indicated by Hoechst staining (fig 5D) without changing the global chromatin accessibility as shown by our ATAC-seq analysis.

Taken together, our transcriptomic and epigenomic wide studies demonstrate that the transcriptional reprogramming of genes involved in mitochondrial function and global translation cannot be assigned to either changes in DNA methylation or chromatin accessibility signatures (RRBS and ATAC-seq) but more likely to the relocation of epigenomic marks and/or a remodeling of the nuclear architecture.

Lmna^{LCS/LCS} pancreatic islets display increased mitochondrial biogenesis and increased global translation.
Since mitochondrial homeostasis is critical for both the maintenance of β-cell function and lifespan promotion (14), we next focused on the regulation of mitochondrial gene expression and function. First, we validated our RNA-seq data by quantitative RT-PCR analysis and confirmed that Lmna<sup>LC5/LCS</sup>-derived islets expressed increased levels of key genes involved in the control of mitochondrial function, including the genes encoding electron transport chain proteins (Nduf, Atp5 and Cox protein family, Fig 6A). In addition, the amount of mitochondrial DNA relative to nuclear DNA was significantly higher in Lmna<sup>LC5/LCS</sup> islets (Fig 6B). Ultrastructural analysis of β-cells by transmission electron microscopy (Fig 6C) together with the oxygen consumption rate (OCR) of islets (Fig 6D) further supported a higher number of functional mitochondria in Lmna<sup>LC5/LCS</sup> islets. Taken together, these results suggest that both mitochondrial number and activity are increased in pancreatic islets isolated from Lmna<sup>LC5/LCS</sup> mice.

RNA-seq data also highlighted a role for lamin C in the regulation of protein translation-related genes in islets from old Lmna<sup>LC5/LCS</sup> mice. In particular, genes involved in ribosome synthesis (Rpl and Mrpl protein family) and in the initiation phase of eukaryotic translation (eIFs, eukaryotic initiation factor) were overexpressed in Lmna<sup>LC5/LCS</sup> islets. qPCR experiments confirmed that the expression of genes directly implicated in different steps of translation was significantly increased in Lmna<sup>LC5/LCS</sup> islets compared to Lmna<sup>+/+</sup> islets (Fig 6E). Because ribosome synthesis is tightly correlated with ribosomal RNA (rRNA) synthesis, we also measured 18S and 28S rRNA levels in isolated islets and found that their expression levels were 1.5-fold increased in Lmna<sup>LC5/LCS</sup> islets (Fig 6F). To confirm that translation rates are increased in Lmna<sup>LC5/LCS</sup> islets, we measured 35S-Met/Cys incorporation and found that global protein synthesis levels were significantly increased in Lmna<sup>LC5/LCS</sup> islets (Fig 6G). Since insulin is the major protein synthetized in β-cells, we measured insulin content in Lmna<sup>+/+</sup> and Lmna<sup>LC5/LCS</sup> isolated islets. Since Lmna<sup>LC5/LCS</sup> islets are larger than control islets, classical insulin content
measurement may not quantitatively reflect the abundance of insulin per cell. Therefore, we performed FACS analysis to measure intracellular insulin levels in dissociated β-cells. As shown in Fig 6H, quantification of FACS experiments revealed that insulin production was increased by approximately 25% in Lmna<sup>LCS/LCS</sup> β-cells.

Altogether, these data suggest that the increased insulin secretion observed in Lmna<sup>LCS/LCS</sup> mice is likely due to an increase in global translation rates, which enhances insulin production. The secretion is supported by an increased number of functional mitochondria that play a crucial role in insulin exocytosis upon glucose stimulation. The observed lamin C-mediated increase in insulin production and secretion may explain the improved metabolic phenotype observed in obese, aged Lmna<sup>LCS/LCS</sup> mice.

**Discussion**

Loss of β-cell function and mass is involved in the development of T1D and T2D, and may contribute to the metabolic complications observed during aging. Our results demonstrate for the first time that the exclusive expression of lamin C acts as a key regulator of β-cell function and regeneration, thus protecting against hyperglycemia observed in several forms of metabolic stresses, such as aging, diet-induced obesity and STZ-induced diabetes. By combining *in vitro*, *ex vivo* and *in vivo* studies, we report that lamin C expression increases β-cell number during aging, counteracts β-cell failure during diet-induced obesity and promotes β-cell regeneration after STZ treatment. Most importantly we demonstrate that lamin C modulates a specific transcriptional program that contributes to enhance the translation of insulin and promotes insulin secretion through a mitochondria-coupled mechanism.

It has been shown that β-cell turnover decreases with aging due to the onset of senescence (15). Indeed, the expression of p16<sup>ink4a</sup>, a key marker of senescence, limits islet
proliferation and regeneration in an age-dependent manner (16). Conversely, the downregulation of p16\textsuperscript{Ink4a} in a \textit{Pten}-null background reduces the age-induced decline of β-cell proliferation and enhances islet mass in aged mice (17), as observed in our aged \textit{Lmna}\textsuperscript{LCS/LCS} mice. While one may speculate that a decrease in cellular senescence in our \textit{Lmna}\textsuperscript{LCS/LCS} mice may underlie the observed maintenance of β-cell mass during aging, no difference in the expression of the senescence marker \textit{p16}\textsuperscript{Ink4a} was observed in our \textit{Lmna}\textsuperscript{LCS/LCS} mice versus controls (data not shown), suggesting that delayed senescence does not contribute to the maintenance of functional islets in our model. This hypothesis is further strengthened by the observation that β-cells which exclusively express lamin C adapt to pathological conditions unrelated to aging, such as diabetes induced in young mice.

The specific functions of lamin A versus lamin C in β-cell maintenance and function during aging and/or metabolic stress is a central question. A limitation of the \textit{Lmna}\textsuperscript{LCS/LCS} model is that we cannot rule out potential contributing effects of lamin C in other organs, as well as an effect of lamin A deficiency in cells expressing lamin C only. Since the gene targeting strategy employed here to generate the \textit{Lmna}\textsuperscript{LCS/LCS} model precludes to specifically express lamin C only in β cells, and subsequently perform lineage-tracing experiments, this question remains to be addressed. Interestingly, mice expressing lamin A only (\textit{Lmna}\textsuperscript{LAO}) are healthy with no detectable phenotypical or histopathological abnormalities, despite a complete loss of lamin C expression (18), even in 24-month-old mice. Challenging these \textit{Lmna}\textsuperscript{LAO} mice with HFD or STZ treatment may help to accurately characterize their specific functions in glucose homeostasis.

To decipher the molecular mechanism by which lamin C regulates pancreatic islet functions, we analyzed the impact of lamin C exclusive expression using genome-wide DNA methylation, ATAC-seq and high-throughput RNA sequencing.
RNA-seq analysis revealed that the expression of genes involved in mitochondrial function and protein translation was altered in aged $Lmna^{LCS/LCS}$ islets compared to controls. Unlike the decline of mitochondrial function observed in aging-associated diseases such as diabetes (14), we found increased mitochondrial activity in aged $Lmna^{LCS/LCS}$ islets that likely contributes to the observed improvement in insulin secretion. It is interesting to note that decreased mitochondrial biogenesis was previously observed in adipose tissue from $Lmna^{LCS/LCS}$ mice aged 45 weeks (6). Contrary to white adipose tissue where mitochondria are relatively less abundant and crucial for adipocyte homeostasis in normal diet and temperature conditions, mitochondrial function in the β-cell is directly coupled to insulin biosynthesis and insulin exocytosis. Since both studies were not conducted in animals at the same ages (45 weeks (6) vs 75 weeks old in the present study), we cannot exclude an age-dependent effect in these tissues. We can however propose that the differential expression of lamin isoforms may have a tissue-specific impact on mitochondrial gene transcription.

RNA-seq analysis and translation rate analysis revealed an increase of global protein synthesis in the islets of aged-$Lmna^{LCS/LCS}$ associated with an increased lifespan as described in our previous study (6). It is widely accepted that reduced protein synthesis promotes longevity (19). Indeed, protein biosynthesis in aging is accompanied by more frequent production of damaged proteins due to transcriptional or translational errors and co-translational protein misfolding. The increased lifespan associated to an upregulation of global protein synthesis in our aged-$Lmna^{LCS/LCS}$ mice suggests that proteins synthetized within the pancreatic islets are well-folded and biologically active, possibly due to the maintenance of an efficient protein quality control system. This hypothesis is supported by our demonstration that insulin, the major protein synthesized by β-cells, is properly secreted and efficient during aging, since aged-$Lmna^{LCS/LCS}$ mice remain glucose tolerant and normoglycemic. This functional insulin
overproduction may contribute to the maintenance of a positive feedback loop of insulin-regulated protein synthesis via the mTOR pathway (20).

How Lmna isoforms are able to modulate gene transcription in β-cells remains to be elucidated. Previous studies have described epigenetic dysregulation in pancreatic islets during T2D, showing differential methylation pattern associated with islet dysfunctions (21) and changes in the open chromatin landscape of human islets (22). Moreover, DNA methylation is a dynamic process in aging β-cells. De novo methylation of proliferation-related genes and demethylation of genes related to β-cell function increases insulin secretion during aging (23). Nevertheless, we did not observe any difference in DNA methylation in Lmna<sup>LCS/LCS</sup> versus wild type pancreatic islets. This finding shows that the selective expression of lamin C does not influence overall DNA methylation in pancreatic islets and points to another regulatory mechanism.

In metazoan nuclei, specific lamina-associated domains (LADs) contribute to the overall spatial organization of the genome and are associated with gene repression (24). Cellular defects in cells from progeria patients include widespread alterations in chromatin organization, such as the loss of heterochromatin domains and changes in epigenetic markers (25). Furthermore, it has been shown that lamin A/C can modulate transcription through the regulation of epigenetic factors (26). Therefore, the expression of only one Lmna isoform may have a crucial impact on the spatial compartmentalization of inactive or active chromosome domains, which in turn may have consequences on transcriptional dysregulation. Surprisingly, we did not observe any global changes in chromatin accessibility through ATAC-seq experiments in our mouse models. Nevertheless, Lmna<sup>LCS/LCS</sup> pancreas show an active enhancer mark (H3K27Ac) relocalization to peripheral heterochromatin. The localization of an active histone mark in an heterochromatin environment has already been described showing that
acetylation of histone in silent domains drive their mislocalization and derepression (27). Furthermore, there is increasing evidence for active and regulated transcription in the heterochromatin at the nuclear periphery (28). Altogether, these observations lead us to speculate that the phenotype observed in \textit{Lmna}^{LCS/LCS} mice could be due to transcriptional reactivation of a subset of mitochondrial and translational genes through histone marks relocalization rather than by reorganizing global chromatin accessibility or modifying DNA methylation. The role of lamin C in the nuclear localization of specific loci and their respective histone marks requires further investigations.

In conclusion, \textit{Lmna}^{LCS/LCS} mice present an enhanced adaptive response to the increased metabolic demands during the course of diabetes and aging. This is probably due to efficient glucose-sensing and insulin production by their neogenerated islets (29). Our study reveals an important role for lamin C in pancreatic \(\beta\)-cells, namely a protective role against metabolic defects arising from \(\beta\)-cell dysfunction. This effect may promote the healthy aging of the \textit{Lmna}^{LCS/LCS} mice, and sustain their extended lifespan (6). The discovery of new drugs targeting \textit{Lmna} alternative splicing may represent an innovative therapeutic strategy to counteract aging \(\beta\)-cell decline and improve diabetes treatment.
Acknowledgements

This work was supported by grants from OSEO-ISI CaReNA (I 13 03 008W to JT), from E.G.I.D (ANR-10-LABX-46 to J-S. A.), ANR (BETAPLASTICITY, ANR-17-CE14-0034 to J-S. A.), EFSD (J-S.A.), I-SITE ULNE (EpiRNA-Diab to J-S. A.), Société Francophone du Diabetes (J-S.A.), INSERM, CNRS, Lille University, Association pour la Recherche sur le Diabète (to J-S.A.), Conseil Régional Hauts de France and Métropole Européenne de Lille (to X.G. and J-S.A.) and from the Swiss National Science Foundation (Ambizione PZ00P3-168077 to ICLM). We thank David Lleres and Cyril Esnault for discussions and scientific critique. The authors are grateful to MRI and RHEM for assistance in cytometry and experimental histology, as well as the animal facility platforms RAM and ZEFI. We especially thank Sylvain de Rossi, Myriam Boyer and Chantal Cazevieille for their technical assistance. We thank Magali A. Ravier for advice for the islet isolation from mice and Amal Makrini for the bioinformamatics support. Marion de Toledo and Carine Chavey are the guarantors of this work.

Authors’ contribution:

CC and MdT designed, performed and interpreted the majority of the experiments. PC performed histological analysis. MP and CB contributed to the RNA-seq and RRBS analysis as well as the bioinformatics analysis. XG gave advice for islet isolation from mice. ICLM contributed to the aging experiments. CC and MdT wrote the manuscript. ICLM, JT and J-SA reviewed and approved the manuscript.

Conflict of interest:

The authors have declared that no conflict of interest exists.
Abbreviations:

AUC: area under the curve, IPGTT: intra peritoneal glucose tolerance test, GSIS: glucose-stimulated insulin secretion, STZ: streptozotocin, T1D: type-1 diabetes, T2D: type-2 diabetes, DIO: diet-induced obesity, HFD: high-fat diet, CD: chow diet, RRBS: Reduced representation bisulfite sequencing.
References


Legends

Figure 1: \textit{Lmna}^{LCS/LCS} mice become obese but remain glucose tolerant as they age.

A) Body weight gain curve in \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (n=10 per group).

B) Fasting blood glucose in young (25 weeks old) and old (75 weeks old) \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (n=5 per group).

C) Intraperitoneal glucose tolerance test (IPGTT) in young and old \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (n=5 per group). The bar graphs (inset) represent the area under the curve (AUC).

D) Fasting insulin levels before and after 30 minutes after intraperitoneal glucose injection in young and old \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (n=5 per group).

All values are expressed as the mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001, n.s not significant by Student’s unpaired t-test.

Figure 2: \textit{Lmna}^{LCS/LCS} mice show increased β-cell mass and insulin secretion during aging

A) Pancreas weight in young (25 weeks old, n=6 per group) and old (75 weeks old, n=17 per group) \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice.

B) Representative images of hematoxylin and eosin (HE) staining of pancreatic sections from young and old \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (scale bar, 2 mm). The right image shows a magnification (4X) of the left image.

C) Scatter plot of the diameters of islets of Langerhans from pancreatic sections of young and old \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (n=5 animals per group).

D) Representative images from immunofluorescence analysis of pancreatic sections showing co-expression of insulin (green) and glucagon (red) in islets from young and old \textit{Lmna}^{+/+} and \textit{Lmna}^{LCS/LCS} mice (scale bar 100 µm).
E) Quantification of β/α cells ratio per pancreatic section in young and old Lmna+/+ and LmnaLCS/LCS mice (n=4)

F) Quantification of isolated islets number after collagenase digestion in young and old Lmna+/+ and LmnaLCS/LCS mice.

G) Global insulin content from total pancreata of old Lmna+/+ and LmnaLCS/LCS mice (n=6 per group)

H) Glucose-stimulated insulin secretion (GSIS) under low (3mM) or high (20mM) glucose concentration from isolated islets from young and old Lmna+/+ and LmnaLCS/LCS mice (n=4 per group). The results were normalized to total insulin content.

All values are expressed as the mean ± SEM. **p < 0.01, ***p < 0.001, n.s not significant by Student’s unpaired t-test.

Figure 3: Increased insulin secretion in LmnaLCS/LCS mice can counteract obesity-induced type 2 diabetes

A) Mean body weight from age-matched Lmna+/+ and LmnaLCS/LCS mice (40 weeks old, n=8 per group) fed chow diet (CD) or a high fat diet (HFD) for 25 weeks.

B) Fasting blood glucose levels over time in Lmna+/+ and LmnaLCS/LCS mice (from 15 to 40 weeks old) during HFD (n=8 per group).

C) Intraperitoneal glucose tolerance test (IPGTT) results in Lmna+/+ and LmnaLCS/LCS mice (40 weeks old) fed a HFD for 25 weeks (n=8 per group). The bar graph (inset) represents the area under the curve (AUC).

D) Mean insulin levels before and 30 min after glucose injection in Lmna+/+ and LmnaLCS/LCS mice (40 weeks old) fed a HFD for 25 weeks (n=8 per group).
E) Mean GSIS under low (3mM) or high (20mM) glucose concentration in isolated islets from 
*Lnma*+/+ and *Lnma*<sub>LC/LCS</sub> mice (40 weeks old) fed a HFD for 25 weeks (n=8 per group). Insulin 
secretion is normalized to total insulin content. 
All values are expressed as the mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001, 
n.s not significant by Student’s unpaired t-test. 

Figure 4: Streptozotocin-injected *Lnma*<sub>LC/LCS</sub> mice display increased β-cell regeneration 
A) Fasting blood glucose levels after streptozotocin (STZ, at day 0) injection in 13-week-old 
*Lnma*+/+ and *Lnma*<sub>LC/LCS</sub> mice (n=5 per group). 
B) Blood glucose levels during IPGTT after 16 days of STZ treatment in *Lnma*+/+ and *Lnma*<sub>LC/LCS</sub> 
mice (15 weeks old, n=5 per group). The bar graph (inset) represents the area under a curve 
(AUC). 
C) Representative immunostaining images of pancreatic sections from *Lnma*+/+ and *Lnma*<sub>LC/LCS</sub> 
mice before and 18 days after STZ injection (n=5 per group) showing co-expression of insulin 
(green) and glucagon (red). Nuclei are counterstained with Hoechst (left panel) (scale bar 100 
µm). Quantification of the mean number of β-cell positive islets/pancreatic section is 
summarized in the bar graph in (right panel). 
D) Representative immunostaining images of pancreatic sections from *Lnma*+/+ and *Lnma*<sub>LC/LCS</sub> 
mice 72 hours after STZ injection (n=5 per group) (left panel) showing the destruction of insulin 
positive b-cells (green) but not glucagon positive a-cells (red). Nuclei are counterstained with 
Hoechst showing a nuclei free area (white dotted line)(scale bar 100 µm). Quantification of the 
β-cells /α-cells ratio before and 72h post-STZ injection is represented in right panel.
E) Representative immunostaining images of pancreatic sections from \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice 9 days after STZ treatment (n=7) showing expression of phospho-histone H3 (red) and insulin (green). Histogramm represents the quantification of the number of phospho-histone H3-positive cells/pancreas section.

F) Representative images of immunostaining of pancreatic sections showing co-expression of specific markers of phospho-histone H3 (red) with either insulin, lectin, glucagon, or somatostatin (top to bottom, green) in \( Lmna^{LCS/LCS} \) mice. Nuclei are stained with Hoechst (scale bar 100 µm).

All values are expressed as the mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001; ns not significant by Student’s unpaired t-test.

Figure 5: \( Lmna^{LCS/LCS} \) mice display a transcriptional reprogramming of genes involved in mitochondria and translation

A) Heatmap showing genes associated with mitochondrial function (upper panel) and translation (lower panel) with differential expression in isolated islets from old \( Lmna^{LCS/LCS} \) versus \( Lmna^{+/+} \) mice (n=9, 3 replicates of 3 mice). The pie chart summarizes the proportion of these classes of genes in the total RNA-seq dataset. The table summarizes the functions of the genes with differential expression in \( Lmna^{LCS/LCS} \) islets as identified by gene ontology (GO).

B) Scatterplot showing the DNA methylation profile analysed by RRBS of \( Lmna^{+/+} \) vs \( Lmna^{LCS/LCS} \) islets (n=2 per condition).

C) Volcano plot of ATAC-seq from \( Lmna^{+/+} \) vs \( Lmna^{LCS/LCS} \) islets. Peaks which P-adj < 0.01 and Log2FC <-1 or >1 are colored in red. Most of the no differential peaks are colored in grey (n=2 per condition).
D) Representative confocal images from immunohistochemistry analysis of pancreatic sections showing co-expression of insulin (red), H3K27Ac, H3K9Me3 (green) and Hoechst staining (grey) in islets from old \( Lmna^{+/+} \) vs \( Lmna^{LCS/LCS} \) mice (scale bar 20 µm). The enlarged cells presented in the insert are indicated by a white arrow.

**Figure 6:** \( Lmna^{LCS/LCS} \) pancreatic islets display increased mitochondrial biogenesis and increased global translation

A) Relative expression levels of mitochondrial genes assessed by RT-qPCR in isolated islets from old \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice (n=8 per group).

B) Quantification of mitochondrial DNA levels by qPCR of \( mt16s \) and \( mtNDS \) gene normalized to genomic DNA in isolated islets from old \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice (n=8 per group).

C) Representative transmission electron microscopy images at x22000 magnification showing a higher number of mitochondria in \( Lmna^{LCS/LCS} \) β-cells relative to control cells (n=3 per group).

D) Mean oxygen consumption rate (OCR) are presented in basal conditions and following sequential injection of glucose, oligomycin, FCCP and rotenone determined with a Seahorse XFe96 flux analyzer in isolated islets from old \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice (n=4 per group).

E) Relative expression levels of translation-related genes assessed by RT-qPCR in isolated islets from old \( Lmna^{+/+} \) versus \( Lmna^{LCS/LCS} \) mice (n=8 per group).

F) Relative 18S and 28S rRNA expression assessed by RT-qPCR in isolated islets from \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice (n=8 per group).

G) Analysis of global translation rates assessed by \( ^{35}S \)-Met/Cys incorporation in isolated islets from \( Lmna^{+/+} \) and \( Lmna^{LCS/LCS} \) mice (n=7 per group). The scatter plot depicts the mean fold change in protein synthesis.
H) Insulin content evaluated by FACS analysis of dissociated islets from old Lmna<sup>+/+</sup> and Lmna<sup>LCS/LCS</sup> mice (insulin/Alexa 647 staining) (n=4 per group)(left panel). The mean insulin of dissociated islets from old Lmna<sup>+/+</sup> and Lmna<sup>LCS/LCS</sup> mice was quantified (right panel).

All values are expressed as the mean ± SEM. *p<0.05, **p<0.01, ***p<0.001, ****p<0.0001 by Student’s unpaired t-test.
A

![Graph showing weight vs. age for Lmna +/+ and Lmna LCS/LCS genotypes.](imageA)

B

![Bar graph showing fasting glucose levels for Young and Old genotypes.](imageB)

C

![Graph showing blood glucose levels over time for Young and Old genotypes.](imageC)

D

![Graph showing insulin levels for Young and Old genotypes.](imageD)
Figure 4:

A) Graph showing changes in fasting glucose levels over time after STZ injection. The graph compares Lmna +/+ and Lmna LCS/LCS genotypes. There are significant differences indicated by asterisks.

B) Graph showing changes in glucose levels over time (in minutes) with Lmna +/+ and Lmna LCS/LCS genotypes. Similar to A, there are significant differences indicated by asterisks.

C) Images and graphs comparing islet numbers before and 18 days after STZ injection. The graphs show a significant increase in β-cell positive islets for Lmna LCS/LCS compared to Lmna +/+.

D) Immunofluorescence images showing insulin, glucagon, and phospho-H3 merged. The graphs show a significant increase in phospho-H3 positive cells for Lmna LCS/LCS compared to Lmna +/+.

E) Images and graphs showing the β-cell to α-cell ratio before and 72h post-STZ injection. The graphs show a significant decrease in the ratio for Lmna LCS/LCS compared to Lmna +/+.

F) Images showing insulin, lectin, glucagon, somatostatin, and phospho-H3 merged. The graphs show a significant increase in phospho-H3 positive cells for Lmna LCS/LCS compared to Lmna +/+. All images are scale bars.
A) Structure of the targeted allele after homologous recombination. The “Lox-STOP-Lox” cassette insertion in intron 10 of LMNA gene induces the exclusive lamin C expression by promoting the use of lamin C polyadenylation site.

B) Expression of lamin A, lamin C and actin expression in the brain, liver, pancreas and muscle of 10-week-old Lmna+/+ and LmnaLCS/LCS mice.

C) Representative HE staining images of pancreatic sections from 121-week-old Lmna+/+ and 141-week-old LmnaLCS/LCS mice. The right image shows a magnification (4X) of the left image.
A) Quantification of number of α-cells per islet section in young vs old $Lmna^{+/+}$ and $Lmna^{LCS/LCS}$ mice (n=4 per group)

B) Glucagon content from total pancreata of old $Lmna^{+/+}$ and $Lmna^{LCS/LCS}$ mice (n=6 per group)

All values are expressed as the mean ± SEM. ns not significant by Student’s unpaired t-test.
A) Lamin A and lamin C expression in CRISPR/Cas9-generated MIN6\textsuperscript{wt}, MIN6\textsuperscript{LCO} and MIN6\textsuperscript{LCO} cells as assessed by western blot.

B) GSIS in MIN6\textsuperscript{wt} and MIN6\textsuperscript{LCO} cells. The results were normalized to insulin content.

All values are expressed as the mean ± SEM. ****p < 0.0001 and ns not significant by Student’s unpaired t-test.
Mean food intake of CD or HFD in 30-week-old *Lmna*<sup>+/+</sup> and *Lmna*<sup>LCS/LCS</sup> mice (n=8 per group).
A) Body weight curve over time in Lmna\(^{+/+}\) and Lmna\(^{LCS/LCS}\) mice during STZ treatment (n=5 per group).

B) Representatives images (left panel) of TUNEL staining (green) on pancreatic sections from Lmna\(^{+/+}\) and Lmna\(^{LCS/LCS}\) mice obtained 48 h after STZ treatment (n=3 per group). Insulin is stained in red, and nuclei are stained with Hoechst (scale bar 50 µm). Quantification (right panel) of the number of TUNEL positive cells per islets.

C) FACS quantification of mean percentages of apoptotic MIN6\(^{wt}\) and MIN6\(^{LCO}\) cells treated with or without STZ (12 mM) for 5 h.

All values are expressed as the mean ± SEM. *p<0.05 and ns not significant by Student’s unpaired t-test.
Distribution of the analyzed CpG sites in various regions of the genome.
Supplementary material

Western Blot

Mouse brain, liver, pancreas and muscle tissue samples were snap-frozen in liquid nitrogen immediately after collection. Samples were then reduced into powder form with a tissue grinder, homogenized and dissolved at 150 mg/ml in a buffer containing 63mM Tris-HCl (pH7.5), 2% (w/v) SDS, 5% (v/v) 2-mercaptoethanol and 8M urea. Samples were stored at -80°C. Protein extracts were size-fractionated on 4-12% gradient SDS-polyacrylamide Bis-Tris gels (ThermoFischer Scientific) and transferred to nitrocellulose membranes for Western blotting. Western blots were probed with a 1:1000 dilution of anti-lamin A/C (sc-20681-Santa Cruz Biotechnology, Santa Cruz, CA, USA), 1:4000 dilution of anti-Actin, clone C4 (#MAB1501 - Millipore, MA, USA).

Food intake

Food intake was measured three times a week over a period of one month with ten animals per genotype in individual cages.

Glucose-stimulated insulin secretion

For insulin secretion, 5 islets of 150–200μm diameter kept in 1,5ml microtubes were pre-incubated for 1 h at 37°C in Krebs–Ringer bicarbonate (KRB) buffer containing: 115mM NaCl, 4.7mM KCl, 6.2mM CaCl2, 20mM NaHCO3, 1.2mM MgSO4, 1.2mM KH2PO4, 16mM HEPES, 0.5% Free Faty Acid BSA, 10mM glucose. The pre-incubation medium was then replaced with KRB buffer supplemented with different glucose concentrations (Low: 3mM; High: 20mM). After incubation for 1 h at 37°C, the supernatant was collected and stored at -
80°C for later analysis of insulin content by ELISA (Mouse Ultrasensitive Insulin elisa-ALPCO, USA). Insulin content is collected after islets lysis in an Acid/Ethanol lysis buffer. Data are expressed as a ratio of insulin secreted/total insulin content.

For MIN6 GSIS assays, prior to glucose stimulation, cells were first starved in KRB buffer without glucose for 1h.

**Total pancreatic insulin and glucagon content**

Mouse pancreata were snap-frozen in liquid nitrogen immediately after collection. Samples were reduced into powder, resuspended in 2ml Acid/Ethanol lysis buffer and incubated O/N at 4°C. After sonication, samples were neutralized with 10% of 1M Na2CO3 and centrifuged. Total Insulin and Glucagon content were measured by ELISA (Mouse Ultrasensitive Insulin elisa-ALPCO, Mercodia Glucagon ELISA - 10 μL).

**Flow cytometry**

The islets were first dissociated into a free cell suspension with 0.05%Trypsine-0.5 mM EDTA. Cells were fixed with 1% PFA in PBS for 10 minutes, washed with PBS and permeabilised with 0.05% Triton X-100 for 10 minutes at 4°C. After blocking with PBS containing 1% BSA, 2% donkey serum, 50 mM glycine, and 0.5% gelatin, cell suspension was incubated with a guinea pig anti-insulin antibody (ab7842; Abcam) for 30 minutes at 4°C, washed and then reacted with a FITC-conjugated donkey anti-guinea pig antibody (F(ab’)2 fragment; Jackson ImmunoResearch Labs). The cell suspension was then washed and analyzed by FACSCANTO II (Becton Dickinson).
**MIN6 cell culture and apoptosis**

The mouse insulinoma MIN6 pancreatic β-cell line (AddexBio) was maintained at 37°C in DMEM High Glucose, Glutamax, and sodium pyruvate (#31966-021, Thermo Fisher Scientific, MA, USA) supplemented with 15% fetal calf serum, 0.05 mM beta-mercaptoethanol and 100 µg/ml penicillin-streptomycin in an atmosphere of 95% air/5% CO₂.

To evaluate the MIN6-induced apoptosis, cells were treated with STZ (12 mM) before staining with Annexin-V-FITC (Invitrogen) and propidium iodide and then analyzed by flow cytometry (FACSCalibur, Becton Dickinson).

**CRISPR-Cas9-mediated LCO MIN6 cell line**

To generate the MIN⁴⁰ cell line, we targeted exon 11 of the *Lmna* gene to enhance the use of the polyA site of lamin C located in intron 10. Gene-specific CRISPR sgRNA sequences were selected to generate a non-homologous end joining in exon 11 of the *Lmna* gene (gene ID: 4000) (Exon11-2119: 5'-ACTCGAAGCTTCCGCAGTGG-3').

The generated sgRNAs were inserted into the SIGMA pU6-gRNA. Both pU6-gRNA and pCMV-Cas9wt-2A-GFP or pCMV-Cas9 D10A-2A-GFP (SIGMA) were cotransfected into MIN6 cells using Lipofectamine 3000 Reagent (Thermo Fisher Scientific). GFP-positive cells were sorted by flow cytometry (FACSCalibur, Becton Dickinson), diluted, reseeded in 100 mm petri dishes and cultured for 2 weeks to obtain isolated clones. After handpicking clones for isolation, cells were collected and analyzed by western blot.

**Histological studies**
For paraffin treatment, pancreases were fixed with 4% paraformaldehyde, dehydrated, embedded in paraffin, sectioned (4µm), and stained with hematoxylin and eosin (HE) following standard procedures for morphological evaluation. HE staining of pancreatic sections was visualized with a Nanozoomer slide scanner (Hamamatsu). Scanned images were analyzed with NDP View software. The mean diameter of the Langerhans islets was determined by measuring the diameter of at least 150 islets per mouse (n=5) using NDP View software.

**Immunohistochemistry and image acquisition**

For immunohistochemistry, pancreas sections were deparaffinized, rehydrated, and incubated for antigen retrieval in citrate buffer at 95°C for 30 min. Nonspecific antibody binding was blocked by incubation with PBS containing 1%BSA, 0.05% Tween20 for 30 min at room temperature. Sections were then incubated in primary antibodies (see table 2) overnight at 4°C. After being washed, primary antibodies were detected using appropriate secondary antibodies. Nuclei were counterstained with Hoechst® 33342 (2µg/ml, ThermoFischer Scientific).

β-cell apoptosis after STZ treatment was determined using a Click-iT™ TUNEL Colorimetric IHC Detection Kit (#C10625, Thermofischer scientific).

Widefield images were acquired using a Metamorph-driven (Molecular Devices, Sunnyvale, CA) Leica DM6000 fluorescence microscope with a 100X HCX Plan 0.7 - 1.4 NA oil objective (NA 1.32, Leica, Melville, NY) and a Coolsnap HQ2 camera (Photometrics, Woburn, MA). Confocal images were acquired using a Confocal Leica SP5-SMDmicroscope and a Leica 63×/1.4 oil HCX PL APO CS objective controlled using the Leica LAS AF software.
Ultrastructural transmission electron microscopy (TEM)

After isolation, 150 islets of Langerhans per mice and per genotype were immersed in a solution of 2.5% glutaraldehyde in Sorensen’s buffer (0.1 M, pH 7.4) overnight at 4 °C. They were rinsed twice in Sorensen’s buffer and post-fixed in a 0.5% osmic acid for 2 hours at room temperature in the dark. Tissues were dehydrated in a graded series of ethanol solutions (30-100%) and embedded in EmBed 812 with an Automated Microwave Tissue Processor for Electronic Microscopy, Leica EM AMW. Thin sections (70 nm; Leica-Reichert Ultracut E) of each block were collected at different levels. These sections were counterstained with uranyl acetate and observed on a Hitachi 7100 transmission electron microscope (TEM) in the Centre de Ressources en Imagerie Cellulaire de Montpellier (France).

RT-qPCR analysis

Total RNA was extracted from isolated islets using TRIzol (Life Technologies, USA). After precipitation in isopropanol, RNA pellets were washed in cold 75% ethanol and resuspended in nuclease-free water. Samples were quantified and evaluated for purity (260-nm/280-nm and 260-nm/230-nm ratio) with a NanoDrop ND-1000 spectrophotometer. Total RNA (1.5 μg) was then converted to cDNA with the Maxima First Strand cDNA Synthesis Kit (#K1642, Thermo Scientific, US). Gene expression was quantified by SYBR green real-time PCR on the LightCycler® 480 System (Roche Applied Science). Reactions were run in triplicate. The mean expression value for the housekeeping genes cyclophillin and TBP was used as an internal control to normalize variability in expression. Primer sequences are provided in Supplemental Table 1.
Mitochondrial DNA quantification

Isolated islets were digested 1h at 55 °C in a lysis buffer containing 100 mM Tris (pH 8), 5 mM EDTA, 0.2% SDS, 200 mM NaCl, and 0.1mg/ml proteinase K for DNA extraction. Quantitative PCR (qPCR) analysis of DNA samples diluted at 1 ng/μl was performed with primers specific for mitochondrial genes (mtND5, mt16s) and normalized to the mean of two nuclear genes (LN11, HK2). qPCR was performed with a LightCycler® 480 and LightCycler® 480 SYBR Green I Master (Roche).

RNA-Seq, RRBS, ATAC-Seq and Data analysis

RNA-Seq

RNA-Seq libraries construction

RNA-Seq libraries were constructed with the Truseq stranded mRNA sample preparation (Low throughput protocol) kit from Illumina. 400ng of total RNA were used for the construction of the libraries. Using poly-T oligo attached magnetic beads, total RNA was purified in order to obtain only poly-A mRNA. Following purification, the mRNA is fragmented into small pieces using divalent cations under elevated temperature. The cleaved RNA fragments are copied into first strand cDNA using SuperScript II reverse transcriptase, Actinomycin D and random hexamer primers. The Second strand cDNA was synthesized by replacing dTTP with dUTP. These cDNA fragments then have the addition of a single 'A' base and subsequent ligation of the adapter. The products are then purified and enriched with 15 cycles of PCR. The final cDNA libraries were validated with a Fragment Analyzer (Advanced Analytical, Ankeny, IA) and quantified using the Kapa Library quantification kit (Kapabiosystems). On 1 sequencing lane of a flowcell V4, and 1 sequencing lane of a flowcell V2, the 6 libraries were pooled in equal proportions, denatured with NaOH.
and diluted to 16 pM before clustering. Cluster formation, primer hybridization and single-end read, 50 cycles sequencing were performed on cBot and HiSeq2500 (Illumina, San Diego, CA) respectively. Image analyses and base calling were performed using the Illumina HiSeq Control Software and the Real-Time Analysis component. Demultiplexing was performed using Illumina’s conversion software (bcl2fastq 2.18). The quality of the raw data was assessed using FastQC from the Babraham Institute and the Illumina software SAV (Sequencing Analysis Viewer). Potential contaminants were monitored with the FastQ Screen software from the Babraham Institute.

**RNA-Seq data analysis**

We aligned RNA-seq reads to the mouse genome (UCSC mm9) with the splice junction mapper TopHat 2.1.1 (1), which used Bowtie 2.2.9 (2). We downloaded gene model annotations from the UCSC database (genes.gtf 31 August 2017) and GeneIDs from the NCBI (gene2refseq.gz, 31 August 2017). Final read alignments having more than 3 mismatches were discarded. We performed gene counting using the union mode of HTSeq-count 0.9.0 (3). Before statistical analysis, genes with less than 15 reads (cumulating all the analyzed samples) were filtered out. Counts were normalized using the Relative Log Expression (RLE) method as implemented in the Bioconductor (4) package edgeR 3.16.5 (5). Differentially expressed (DE) genes were identified using 3 different statistical methods: DESeq 1.26.0 (6), EdgeR, and DESeq2 1.14.1 (7). We corrected P-values for multiple testing using the Benjamini-Hochberg FDR method (8). Genes with adjusted P-value <0.05 were declared ‘differentially expressed’. The final lists of DE genes were the genes common to the 3 tests. To perform the functional analysis of the resulting list of genes with the Gene Ontology (GO) annotations, the topGO(9) package from Bioconductor was used. Overrepresented GO terms
were identified using Fisher’s exact test with the weight method that is implemented in the topGO package. As confidence threshold we used a P-value of 1%. To perform this analysis the differentially expressed genes were compared with those of all known genes present in the annotation. The GO categories were found in the Org.Mm.eg.db package based on the gene reporter EntrezGeneID.

**Reduced Representation Bisulfite Sequencing**

*Libraries construction*

Library construction was performed using the Premium RRBS kit (Diagenode). Briefly, for each sample, 100ng of intact genomic DNA were digested using MspI for 12 hours at 37°C. DNA ends were repaired and Diagenode indexed adaptors were ligated to each end of the repaired DNA. Each ligated DNA was quantify by qPCR using the Kapa Library quantification kit (Kapabiosystems) on a LightCycler 480 (Roche Life Science) prior to pooling (8 samples per pool). Each pool was submitted to bisulfite conversion and desalted. Optimal PCR cycle number was determined by qPCR using the Kapa Library quantification kit (Kapabiosystems) before the final enrichment PCR. Once purified using magnetic beads (AMPure XP, Beckman Coulter), library pools were verified on Fragment Analyzer (AATI) and precisely quantified by qPCR using the Kapa Library quantification kit (Kapabiosystems). Each pool was denatured, diluted to a 18 pM concentration and spiked with a 10 % phiX Illumina’s library before clustering. Clustering and sequencing were performed in paired end 2*50nt, 2 lanes per pool, according to the manufacturer’s instructions on a Hiseq2500 using Rapid V2 clustering and SBS reagents.

*Analysis of RRBS data*
The sequencing quality control of RRBS data was identical to the one performed for RNA-Seq data. The un-methylated cytosine added at the end-repair step during library preparation was used as a built-in bisulfite conversion efficiency control. Sequences that contained the adapter on their 3’ end were trimmed with cutadapt (https://cutadapt.readthedocs.org/en/stable/; minimal overlap length between the read and the adapter of 5; discard trimmed reads shorter than 15). The option "--trimmed-only" was used to discard reads that did not contain the adapter. The percentage of non-bisulfite converted cytosines was then calculated as the number of times the filled in position at the 2nd base from the 3’ end was not converted to T. Mean ± sd bisulfite conversion was 99.55 ± 0.03% for the read 1 and 99.63 ± 0.02% for the read 2 for the 24 samples.

Trim Galore (http://www.bioinformatics.babraham.ac.uk/projects/trim_galore/) was used to perform adapter trimming and quality control in two subsequent steps in order to remove the 2 additional bases that contained a cytosine which were artificially introduced at the end-repair step during the library preparation (--rrbs option). The mm9 mouse reference genome was prepared for Bismark 0.17.0 using bismark_genome_preparation script. Bismark software (version 0.17.0) from the Babraham Institute (10) was used to align trimmed RRBS reads to the mouse genome. Then, bismark_methylation_extractor extracted the methylation call for every single C analyzed.

Differential methylation analysis was performed using the methylSig (v0.4.4) Bioconductor R package (11). A site specific analysis was done for each CpG site covered by min 10 and max 500 reads using local information to improve variance estimation. A tiled analysis was also conducted by using the total number of cytosines in a windows of 25 bp. A cytosine or a region was declared differentially methylated if the adjusted p-value (according
to Benjamini-Hochberg multiple testing correction) was less than 0.05 and the difference of methylation percent above 25%.

**Assay for Transposase-Accessible Chromatin (ATAC-Seq)**

Cells were harvested and frozen in culture media containing 50% FCS, 40% RPMI and 10% DMSO. Cryopreserved cells were sent to Active Motif to perform the ATAC-seq assay. The cells were then thawed in a 37°C water bath, pelleted, washed with cold PBS, and tagmented as previously described (12), with some modifications based on (13). Briefly, cell pellets were resuspended in lysis buffer, pelleted, and tagmented using the enzyme and buffer provided in the Nextera Library Prep Kit (Illumina). Tagmented DNA was then purified using the MinElute PCR purification kit (Qiagen), amplified with 10 cycles of PCR, and purified using Agencourt AMPure SPRI beads (Beckman Coulter). Resulting material was quantified using the KAPA Library Quantification Kit for Illumina platforms (KAPA Biosystems), and sequenced with PE42 sequencing on the NextSeq 500 sequencer (Illumina).

Analysis of ATAC-seq data was very similar to the analysis of ChIP-Seq data. Reads were aligned using the BWA algorithm (mem mode; default settings). Duplicate reads were removed, only reads mapping as matched pairs and only uniquely mapped reads (mapping quality >= 1) were used for further analysis. Alignments were extended in silico at their 3’-ends to a length of 200 bp and assigned to 32-nt bins along the genome. The resulting histograms (genomic “signal maps”) were stored in bigWig files. Peaks were identified using the MACS 2.1.0 algorithm at a cutoff of p-value 1e-7, without control file, and with the –nomodel option. Peaks that were on the ENCODE blacklist of known false ChIP-Seq peaks were removed. Signal maps and peak locations were used as input data to Active Motifs
proprietary analysis program, which creates Excel tables containing detailed information on sample comparison, peak metrics, peak locations and gene annotations.

For differential analysis, reads were counted in all merged peak regions (using Subread, v1.5.2), and the replicate groups for each condition were compared using DESeq2 (v1.24.0).
References


### RT qPCR primers

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<tr>
<th>Gene</th>
<th>Forward Primer</th>
<th>Reverse Primer</th>
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<tbody>
<tr>
<td>Cyclophillin</td>
<td>5'-GTT GGC CAG GCT GGT GTC CAG-3'</td>
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<tr>
<td>TBP</td>
<td>5'-GTT GGC GGG TAT CTG CTG GC-3'</td>
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<td>Mm Ndufa6</td>
<td>5'-GAA TGC CCA TGT CAC AGA-3'</td>
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<td>mNdufc1</td>
<td>5'-TTC CTC TGG ATT CAC CCT CA-3'</td>
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<tr>
<td>Mm Atp5f1</td>
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<td>Mm Cox16</td>
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<td>Mm Eif1</td>
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<td>Mm Mrpl52</td>
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<td>18S</td>
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### Mitochondrial DNA

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<td>LN11</td>
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<td>mHK2</td>
<td>5'-GCC AGC CTC TCC TGA TTT TAG-3'</td>
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**Supplemental table 1**: List of primers for real-time PCR
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<tr>
<td>Rabbit anti-Phospho-histone H3 Ser10</td>
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<td>Rabbit anti-H3K9Me3</td>
<td>Ab-8898</td>
<td>1/400</td>
<td>AbCam</td>
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<td>Fluorescein-DBA (Dolichos biflorus Agglutinin)</td>
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<td>Goat anti-rabbit IgG Alexa555</td>
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**Supplemental table 2**: List of antibodies