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Melatonin induces fat browning by transdifferentiation of white adipocytes and *de novo* differentiation of mesenchymal stem cells†

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The role of melatonin in obesity control is extensively accepted, but its mechanism of action is still unclear. Previously we demonstrated that chronic oral melatonin acts as a brown-fat inducer, driving subcutaneous white adipose tissue (sWAT) into a brown-fat-like function (beige) in obese diabetic rats. However, immunofluorescence characterization of beige depots in sWAT and whether melatonin is a beige-fat inducer by *de novo* differentiation and/or transdifferentiation of white adipocytes are still undefined. Lean (ZL) and diabetic fatty (ZDF) Zucker rats were subdivided into two groups, control (C) and oral melatonin-supplemented (M, 10 mg kg⁻¹ day⁻¹) for 6 weeks. Mesenchymal stem cells (MSCs) were isolated from both rat inguinal fat and human lipoaspirates followed by adipogenesis assays with or without melatonin (50 nM for 12 h in a 24 h period, 12 h+/12 h-) mimicking the light/dark cycle. Immunofluorescence and western-blot assays showed the partial transdifferentiation of white adipocytes in both ZL and ZDF rats, with increasing thermogenic and beige markers, UCP1 and CITED1 and decreasing white adipocyte marker ASC-1 expression. In addition, melatonin increased UCP1, CITED1, and PGC1- α expression in differentiated adipocytes in both rats and humans. These results demonstrate that melatonin increases brown fat in obese diabetic rats by both adipocyte transdifferentiation and *de novo* differentiation. Furthermore, it promotes beige MSC adipogenesis in humans. This may contribute to the control of body weight attributed to melatonin and its metabolic benefits in human diabetes.

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1. Introduction

Obesity is the result of a complex interaction between genetics, lifestyle and environment, including social and cultural factors.¹ In addition, this disease carries other comorbidities that include type 2 diabetes mellitus (DM2) among others.² Obesity and its complications are also associated with meta-inflammation, mitochondrial dysfunction, cellular apoptosis,

and oxidative stress.² The modern lifestyle enhances obesity and overweight through a large energy imbalance caused by a higher caloric intake with lack of physical activity.³ The latest studies from the World Health Organization revealed that the global prevalence of obesity has tripled between 1975 and 2016.⁴ In 2016, more than 1.9 billion adults aged 18 years and over were overweight (39%; body mass index ≥ 25 kg m⁻²) and 650 million of them were obese (13%; body mass index ≥ 30 kg m⁻²). In the same year, 340 million children and adolescents between the ages of 5 and 19, and another 41 million under the age of 5, were obese or overweight, also raising a growing concern about childhood obesity. Obesity in humans and other mammals leads to the storage of excess energy as triglycerides in adipose tissue. While white adipose tissue (WAT) functions as a lipid storage reservoir, brown adipose tissue (BAT) provides a mechanism for thermogenesis, dissipating energy in the form of heat.⁵ Brown-colored adipocytes have also been found within WAT. They are called beige or brite (brown on white) adipocytes. Interestingly, these beige/brite cells resemble white adipocytes in the basal state, but respond to thermogenic stimuli with higher levels of thermogenic

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genes and higher respiration rates. Furthermore, beige/brite cells have a different gene expression pattern than white or brown fat cells.⁶ While the alanine-serine-cysteine transporter-1 (ASC-1) is highly expressed in WAT with little or no expression in thermogenic adipose tissues such as BAT, suggesting that this transporter is a specific surface marker for WAT,^{7,8} the Cbp/The transactivator that interacts with p300 with the Glu/Asp-rich carboxy-terminal domain 1 (CITED1) is a marker selectively expressed in beige adipocytes of uncoupling protein 1 (UCP1) positive in mice, as well as in humans.^{9–12}

Melatonin is a hormone produced at night by the pineal gland^{13,14} and also in many other tissues.^{15,16} In addition to physiological circadian rhythm regulation, melatonin has antioxidant, anti-inflammatory, and anti-obesogenic effects. Melatonin is a powerful antioxidant, and its presence in numerous plants and foods has been described as beneficial for human and animal health.^{17,18} Melatonin limits obesity in rodents without influencing food intake^{19–23} with inconsistent changes in locomotor activity, either increase,^{19,23} decrease²⁴ or no changes.²¹ Moreover, in this same rat strain, we have shown that melatonin does not affect locomotor activity.²⁵ Furthermore, in this same rat strain, we have shown that chronic melatonin supplementation has an antidiabetic effect accompanied by an improvement of insulin resistance, as evidenced by a lower HOMA-IR index²⁶ and increases systemic levels of adiponectin which improves insulin sensitivity by increasing energy expenditure and fatty acid oxidation through the activation of AMP activated protein kinase (AMPK) and by increasing the expression of PPAR γ target genes²⁷ like PGC1 α , the master regulator of the thermogenic program and mitochondrial biogenesis, and UCP1 its molecular effector.²⁸ This suggests that melatonin may act as a thermogenic agent. In fact, many previous reports indicated that melatonin enhances the growth and function of thermogenic adipose tissue.²⁹ In addition, white subcutaneous fat contains patches of beige adipocytes, in both animals and humans.^{9,30} These fat cells have structural and functional properties similar to brown adipocytes.^{31,32} Beige deposits can be induced in animals and humans by physiological stimuli, such as cold exposure, which increases adrenergic tone,^{31,33} and by physical activity through irisin, a myokine recently identified in animals and humans.³² This discovery is of utmost interest since the search for safe and effective thermogenic agents is a promising strategy to fight obesity. Therefore, melatonin, besides stimulating the growth and thermogenic activity of thermogenic adipose tissue, could have an “irisin-like” effect by inducing the appearance of “brown-like” adipocytes in areas of WAT. In this sense, a study of our group has shown that the chronic oral administration of melatonin induces the appearance of “brown-like” adipocyte nests in the subcutaneous inguinal WAT from ZDF rats.²⁵ Moreover, we have observed a basal inguinal thermogenic activity and a cold-induced sensitizing regional action in response to the systemic administration of melatonin. The current epidemic of obesity has increased the interest in studying adipocyte formation (adipogenesis), especially in beige/brite adipocytes, as they could shed light on

the search for new effective treatments.^{34,35} Numerous lineage experiments provide evidence that adrenergic stimulation and/or cold exposure increase beige adipocyte biogenesis (browning) in WAT by *de novo* differentiation of pre-adipocyte cells found in WAT depots (adipogenesis), and/or reversible trans-differentiation of mature white adipocytes whose commitment reversibly changes to a beige phenotype, leading to the direct differentiation of mature white to beige adipocytes.^{6,36–44} Several studies also revealed that melatonin and other hormonal factors like triiodothyronine (T3), parathyroid hormone (PTH), glucagon-like peptide 1 (GLP1), leptin and natriuretic peptides also induce the browning of WAT and/or BAT thermogenesis stimulation.⁴⁵ A recent transcriptional and epigenomic study showed specific expression patterns in beige adipocytes from mice, suggesting a distinct lineage and an epigenomic memory of prior cold exposure for beige adipocytes.⁴⁶ These pieces of evidence may support the hypothesis that beige adipocytes contribute to adipose tissue plasticity and adjusting their gene expression patterns and morphology, depending on the presence and frequency of different stimuli,^{39,41} for a rapid thermogenic response.⁴⁶

In the present study, we investigated the characterization of beige depots in sWAT in ZDF rats and the elucidation of whether melatonin induces beige adipocytes by the differentiation of ZDF rat and human subcutaneous WAT-derived mesenchymal stem cells (MSC) and/or the trans-differentiation of white adipocytes to beige ones in ZDF inguinal fat.

2. Materials and methods

2.1 Animals and experimental protocols

This study was carried out in accordance with the European Union guidelines for animal care and protection and the Ethical Committee of the University of Granada (Granada, Spain). The permit project number is 4-09-2016-CEEA. Male ZDF rats (fa/fa $n = 16$) and male lean littermates (ZL, fa/- $n = 16$) were obtained at the age of 5 weeks. Animals were maintained on Purina 5008 rat chow (protein 23%, fat 6.5%, carbohydrates 58.5%, fiber 4%, and ash 8%; Charles River, France) and housed 2 per clear plastic cage in a climate-controlled room at 28–30 °C and 30–40% relative humidity, with a 12 h dark/light cycle (lights on at 7 a.m.). In the first week after arrival, the animals were acclimated to room conditions, and the water intake was recorded. Then, both ZL and ZDF rats were subdivided into four groups. Animals were supplemented for 6 weeks either with melatonin in drinking water (melatonin-supplemented, M-ZDF, $n = 8$ and M-ZL, $n = 8$) or the vehicle (controls, C-ZDF, $n = 8$ and C-ZL, $n = 8$). Melatonin was dissolved in a minimum volume of absolute ethanol and diluted in drinking water to yield a dose of 10 mg per kg body weight (BW) per day, with a final concentration of 0.066% (w/v) ethanol. Water intake and BW were recorded twice weekly. Fresh melatonin and vehicle solutions were prepared twice a week, and the melatonin dose was adjusted to the BW throughout the study period. Water bottles were covered with



aluminum foil to protect from light. At the end of the supplementation period, the animals were anesthetized with sodium thiobarbital (thiopental) and sacrificed.

2.2 Immunofluorescence

Inguinal adipose tissue from Zucker rats was fixed by immersion in 10% formaldehyde for 12 h and after dehydration by immersion in increasing concentrations of ethanol (70–100%), they were included in paraffin forming blocks. Paraffin-embedded inguinal adipose tissue from Zucker rats was cut into 7 μm sections. Before staining, the cuts were deparaffinized with xylene and decreased concentrations of ethanol (100–70% and water). They were subjected to citrate-based antigen retrieval using Antigen Retrieval Citra Solution (Biogenex) and blocked for 1 h at room temperature with rabbit polyclonal antibodies. The sections were incubated for 1 h at room temperature with a mouse monoclonal antibody against CITED1 at a dilution of 1:1000 (Sigma, Spain) or a rabbit monoclonal antibody against UCP1 at a dilution of 1:1000 (Sigma, Spain) or immunoglobulin G (IgG) as a negative control. Secondary detection was performed for 1 h at room temperature using anti-mouse Fluorescein isothiocyanate (FITC) and anti-rabbit Cyanine 3 (Cy3) (Invitrogen, Carlsbad, CA, USA). The cuts were labeled with 4',6-diamidino-2-phenylindole (DAPI). Images were captured using an Olympus IX2 microscopy. Image analysis and quantification was carried out by measuring signal intensities using the ImageJ programmer.

2.3 Isolation of mesenchymal stem cells

Adipose tissue samples (~300 mg) from Zucker rats were excised from brown-like (beige) and white fat regions of the inguinal fat pad and washed in PBS with 1% (w/v) antibiotics (penicillin-streptomycin (P/S), Sigma, Spain). MSCs from beige (bMSCs) and white (WMSCs) inguinal adipose tissue were isolated by enzymatic digestion (Collagenase type I, Sigma, Spain) at 37 °C for 1 h. The enzymatic solution was inactivated by adding complete Dulbecco's Modified Eagle's Medium (DMEM, Gibco, Life Technologies, Spain) supplemented with 10% Fetal Bovine Serum (FBS) and 1% P/S. The mixture was centrifuged at 400 G for 10 min and the pellet was resuspended in growth medium (complete DMEM supplemented with L-glutamine 2 mM, 10% FBS, 1% P/S and 1% doxycycline).

Human MSCs were isolated from lipoaspirates' adipose tissue. Human abdominal fat was obtained from healthy donors ($n = 8$) undergoing liposuction plastic surgery (range of age 44–61). All samples used in this study were collected with informed consent and Institutional Review Board approval (ethic permission number: 02/022010 Hospital Virgen de la Victoria, Málaga, Spain). Briefly, lipoaspirates were washed and treated for 1 h at 37 °C with 1% collagenase type IA (Sigma Aldrich, Spain) in Hank's Balanced Salt Solution (Sigma Aldrich, Spain) under gentle agitation. The digested tissue was centrifuged, and the supernatant discarded. The cell pellet obtained was washed twice in DMEM supplemented

with 10% FBS and resuspended in growth medium. Approximately 1.8×10^5 (range $1.6\text{--}2 \times 10^5$) cells were obtained from 1 g of lipoaspirate.

2.4 Cell culture

MSCs from inguinal adipose tissue of Zucker rats and human lipoaspirates were seeded and expanded in the appropriate sterile plastic flask at equal densities (5×10^3 cells per cm^2) in T-75 culture flasks with 10 ml of complete culture medium consisting of DMEM (Gibco, Life Technologies, Spain) supplemented with L-glutamine (2 mM), 10% FBS (Gibco, Life Technologies, Spain), 1% P/S (Gibco, Life Technologies, Spain) and 1% doxycycline (Sigma, Spain) and cultured in an incubator at 37 °C with a humidified atmosphere containing 21% O_2 and 5% CO_2 . The cell culture medium was replaced twice a week. Freshly isolated cells were grown in monolayer culture up to passage 4–5 at a seeding density of 5×10^3 cells per cm^2 at each passage.

2.5 MSC characterization

After two passages, MSC specific surface markers were analyzed by flow cytometry with non-conjugated antibodies and *in vitro* differentiation assay to be characterized. Mouse anti-rat/human CD-73 (1:100; BD, Pharmingen, San Jose, CA, USA), mouse anti-rat/human CD-90 (1:100; BD, Pharmingen, San Jose, CA, USA), mouse anti-rat/human CD-105 (1:100; BD, Pharmingen, San Jose, CA, USA), mouse anti-rat/human CD-29 (1:100; BD, Pharmingen, San Jose, CA, USA), mouse anti-rat anti-SCA-1 (1:100; BD, Pharmingen, San Jose, CA, USA), mouse anti-rat CD-133 (1:100; BD, Pharmingen, San Jose, CA, USA), mouse anti-rat/human CD-45 (1:100; Millipore, Burlington, MA, USA) and mouse anti-rat/human CD-34 (1:100; BD, Pharmingen, San Jose, CA, USA) were used. As a secondary antibody, bovine anti-mouse FITC (fluorescein isothiocyanate, green fluorescence, Santa Cruz Biotechnology Inc., Santa Cruz, CA, USA) was used for all samples. Isotype control could not be carried out because all the primary antibodies were produced in mice. As negative controls, cell aliquots were incubated only with isotype-matched mouse IgG conjugated to FITC (Santa Cruz Biotechnology Inc., Santa Cruz, CA, USA) under the same conditions. To carry out this procedure, cultured cells in a T-175 flask were washed with PBS and detached by incubation with trypsin diluted 1:1 in PBS. After trypsin inactivation with complete medium and centrifugation at 1500 rpm for 5 min, pellets were resuspended with 10 ml of Blocking Buffer, which contains 2–4% of bovine serum albumin (BSA) and 2 μM ethylenediaminetetraacetic acid (EDTA) in PBS. During 1 h incubation at 37 °C, this solution blocked nonspecific binding between the antibody and other regions. Then, 6 ml of the resuspension was distributed into 6 cytometer tubes, 1 ml in each one. The samples were centrifuged at 1500 rpm for 5 min and the pellet was resuspended with 90 μl of PBS and 10 μl of each primary antibody, for the two cell lines, not including the controls. After incubation at 37 °C for 15–20 min, the samples were washed with 5 ml of PBS by centrifugation at 1500 rpm for 5 min to elimin-



ate the antibody excess and reduce false positives and background signal. 10 μ l of the secondary antibody and 90 μ l of PBS were added, to each sample, which were then incubated at 37 °C for 20 min under dark conditions. After that, the samples were washed with 5 ml of PBS by centrifugation at 1500 rpm for 5 min before being analyzed using a FACScan flow cytometer (BD Biosciences, San Jose, CA, USA).

For the MSC multipotent differentiation potential characterization, *in vitro* adipogenic, osteogenic, and chondrogenic differentiation assays were performed. MSCs were plated at the density of 300 000 cells per well in a three 6-well plate one for each differentiation assay with adipogenic (DMEM, 10% FBS, 1% P/S, 1 mM dexamethasone, 500 μ M isobutyl methylxanthine (IBMX), 60 μ M indomethacin and 10 μ g ml⁻¹ insulin), osteogenic (DMEM, 10% FBS, 1% P/S, 10 nM dexamethasone, 5 mM β -glycerophosphate and 50 μ g ml⁻¹ ascorbic acid) and chondrogenic (DMEM, 10% FBS, 1% P/S, 1% insulin-transferrin-selenium, 100 nM dexamethasone, 40 μ g ml⁻¹ proline, 10 ng ml⁻¹ transforming growth factor-beta 3 (TGF β 3) and 50 μ g ml⁻¹ ascorbic acid) medium, respectively. Cells were cultured for 21 days at 37 °C in a humidified atmosphere containing 21% O₂ and 5% CO₂, changing the medium twice a week to perform complete MSC differentiation into adipocytes, osteocytes, and chondrocytes. After the *in vitro* differentiation assay, differentiated cells and MSCs were fixed with 4% paraformaldehyde (30 min at room temperature), washed twice with PBS, stained with Oil Red O (adipocytes), Alizarin Red (osteocytes) or toluidine (chondrocytes) (30 min at room temperature) followed by three washes with PBS, and visualized with an optical microscope and photographed.

2.6 *In vitro* melatonin treatment

ZDF rat and human MSCs were subjected to an *in vitro* melatonin treatment to differentiate them into adipocytes. MSCs were plated at the density of 30 000 cells per cm² in a 12-well plate with pre-adipocyte growth medium (PGM) consisting of DMEM, 10% FBS and L-glutamine (2 mM) and grown to confluence for 2 days. Cells were then incubated for 10 days in a pre-adipocyte differentiation medium (PDM), to achieve the commitment of cells to the adipogenesis, with or without melatonin. Melatonin was used at 50 nM and was added intermittently in melatonin-treated samples (for 12 h in a 24 h period, 12 h+/12 h-, mimicking the light/dark cycle). The PDM consisted of PGM with 1 μ M dexamethasone (Sigma, St Louis, MO, USA), 60 μ M indomethacin (Sigma, Spain), 500 μ M 3-isobutyl-1-methylxanthine (IBMX, Sigma, Spain) and 0.01 mg ml⁻¹ insulin (Sigma, Spain). Then, the PDM was replaced with an adipogenic differentiation medium (ADM, Lonza, Basel, Switzerland) and cells were incubated 10 days more. Adipogenesis was monitored by the morphological examination of the cellular accumulation of lipid droplets by AdipoRed Assay Reagent (Lonza, Basel, Switzerland) following the manufacturer's protocol. The stained lipid droplets were visualized with an optical microscope and photographed. To quantify adipogenesis, the absorbance of the stained cultures was measured at 520 nm using a microplate reader.

2.7 Quantification of UCP1, ASC-1, PGC1 α and CITED1

ZDF rat and human MSCs and adipocytes obtained after *in vitro* melatonin treatment were harvested in 10 mM Tris-HCL (pH 7.5), 150 mM NaCl, 2 mM EDTA, 1% Triton X-100, 10% glycerol (all from Sigma, Spain) and protease inhibitor cocktail (Thermo Fisher Scientific, Waltham, MA, USA), and placed on ice for 20 min. Following centrifugation (13 000g, 4 °C, 30 min), the protein content of the supernatant was determined using Protein Assay Kit II (BioRad, Spain). For inguinal adipose tissue samples from Zucker rats, about 100 mg were homogenized in lysis buffer (150 mM NaCl, 5 mM EDTA, 50 mM Tris-HCL, pH 7.4) without Triton X-100 and homogenized with a Teflon pestle. The homogenates were centrifuged (3000g, 4 °C, 15 min), and the upper-fat layer was removed from the supernatant. Then, Triton X-100 was added to a final concentration of 1%. After incubating at 4 °C for 30 minutes, extracts were cleared by centrifugation at 15 000g at 4 °C for 15 min and the protein content of the supernatant was determined using Protein Assay Kit II. 100 μ g of total protein of each sample was analyzed by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). The gels for immunoblot analyses were transferred to a nitrocellulose membrane (BioRad Trans-Blot SD; BioRad Laboratories, Hercules, CA, USA). Blots were reacted with a 1 : 2000 dilution of primary antibodies (anti-UCP1, anti-ASC-1, anti-PGC1 α and anti-CITED1, all produced in rabbit, Sigma, Spain) in blocking solution (PBS, 5% BSA). The anti- β -actin antibody generated in mice (Santa Cruz Biotechnology, Santa Cruz, CA, USA) was used as a control. After overnight incubation at 4 °C with the primary antibody, horseradish peroxidase-conjugated anti-rabbit and anti-mouse secondary antibody incubation was performed. Immune reactive signals were detected by enhanced chemiluminescence using a Clarity Western ECL Substrate kit (BioRad, Spain). Blots were digitally recorded with Kodak Image Station 4000 PRO and analyzed by image analysis for molecular weight determinations. Fold changes in relative expression were calculated from the signal intensities derived using the ImageJ programmer.

2.8 Statistical analysis

All experiments were repeated at least three times. Data are expressed as means \pm SEM. Means were compared between groups using one way-ANOVA, adjusted by Tukey. SPSS, version 22, for Windows (SPSS, Michigan, IL, USA) was used for the data analyses. *P*-Value < 0.05 was considered statistically significant, and the levels of significance were labeled on the figures as follows: ***P* < 0.01; **P* < 0.05; and ###*P* < 0.01; #*P* < 0.05.

3. Results

3.1 *In vivo* melatonin supplementation induces partial white to beige adipocyte transdifferentiation in the inguinal WAT of obese diabetic (ZDF) rat

In order to immunocharacterize beige depots, we carried out an immunofluorescence study of inguinal WAT on ZDF and ZL



rats, supplemented either with melatonin (M-ZDF and M-ZL) or the vehicle (controls, C-ZDF and C-ZL). DAPI blue fluorescence (Fig. 1A) showed a decreased number of cells in the inguinal WAT from ZDF rats compared to lean littermates, with a higher relative cell number in the M-ZDF group compared to the C-ZDF one (C-ZL, 1.046 ± 0.066 ; C-ZDF, $0.102 \pm$

0.019 ; M-ZDF, 0.488 ± 0.044 ; $P < 0.01$; Fig. 1B). Immunofluorescence assay showed a significant decrease in the fluorescence signal of all measured markers in the C-ZDF group which is reversed by *in vivo* melatonin supplementation in ZDF rats (Fig. 1C and E). Also, no differences between C-ZL and M-ZL groups were observed in none of the markers deter-

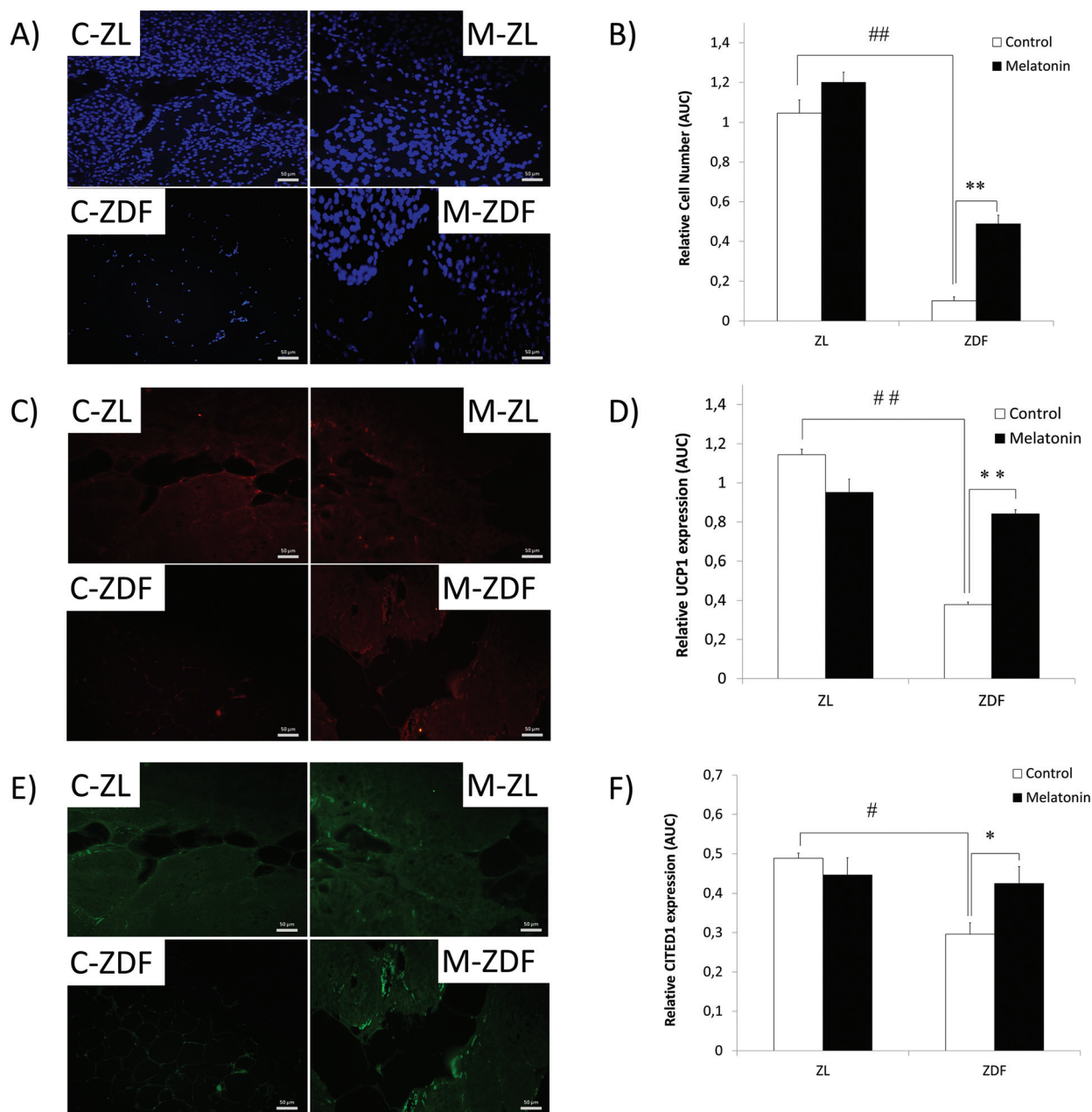


Fig. 1 Immunofluorescence characterization of the “brown-like” cell nests of the melatonin-induced inguinal WAT of the Zucker rats. (A) Representative microscopy image showing blue labeled nucleus (DAPI); (B) quantitative analysis of relative nucleus number corresponding to adipocyte cell number quantification; (C) representative microscopy image showing immunofluorescence in red corresponding to anti-rabbit-monoclonal-UCP1 (Cy3) thermogenic marker; (D) quantitative analysis of the relative expression of thermogenic cell marker UCP1; (E) representative microscopy image showing immunofluorescence in green corresponding to the anti-mouse-monoclonal-CITED1 (FITC) specific beige-cell marker; and (F) quantitative analysis of the relative expression of specific beige-cell marker CITED1. ZL, Zucker lean rats; ZDF, Zucker diabetic fatty rats; C, control-unsupplemented; M, melatonin-supplemented. Magnification 20X. Scale bar 50 μm . All values are expressed as mean \pm SEM of four independent experiments in triplicate. $**P < 0.01$; $*P < 0.05$; $###P < 0.01$; $\#P < 0.05$.



mined by immunofluorescence. *In vivo* melatonin supplementation induces higher UCP1 expression in the inguinal WAT from ZDF rats (C-ZDF, 0.378 ± 0.013 ; M-ZDF, 0.842 ± 0.020 ; $P < 0.01$; Fig. 1D). Furthermore, unsupplemented C-ZDF rats slightly expressed UCP1 and not C-ZL the UCP1 expression of which was higher (C-ZDF, 0.378 ± 0.013 ; C-ZL, 1.144 ± 0.028 ; $P < 0.01$; Fig. 1D). CITED1 expression in unsupplemented obese rats was lower than that in unsupplemented lean littermates too (C-ZDF, 0.296 ± 0.029 ; C-ZL, 0.489 ± 0.013 ; $P < 0.05$; Fig. 1F). *In vivo* melatonin supplementation also significantly induces CITED1 expression in the inguinal WAT from ZDF rats compared to the corresponding control (C-ZDF, 0.296 ± 0.029 ; M-ZDF, 0.425 ± 0.042 ; $P < 0.05$; Fig. 1F). No differences between M-ZDF and C-ZL groups were found, recovering the decreased expression of CITED1 observed in C-ZDF rats. Moreover, to evaluate whether melatonin induces the trans-differentiation of white to beige adipocytes and/or differentiation of MSCs to mature beige adipocytes, white adipocyte quantification in inguinal WAT, UCP1 expression study in MSCs from *in vivo* studies of both lean and obese non-supplemented and melatonin-supplemented rats, and an *in vitro* melatonin treatment study of MSCs from the inguinal fat of Zucker rats were accomplished.

For white adipocyte quantification, the expression of ASC-1, a specific cell surface marker, was measured. Western blot analysis revealed that *in vivo* melatonin supplementation decreases the expression of ASC-1 both in lean (C-ZL, 2.276 ± 0.011 ; M-ZL, 1.488 ± 0.041 ; $P < 0.01$) and diabetic fatty rats (C-ZDF, 2.702 ± 0.079 ; M-ZDF, 2.025 ± 0.026 ; $P < 0.01$) as shown in Fig. 2. These results could explain the origin of mela-

tonin-induced beige adipocytes by partial transdifferentiation of white to beige adipocytes. However, more studies are needed to understand this biological process. Also, ASC-1 expression in non-supplemented obese rats is significantly higher than that in control lean ones (C-ZL, 2.276 ± 0.011 ; C-ZDF, 2.702 ± 0.079 ; $P < 0.05$; Fig. 2).

3.2 *In vivo* melatonin supplementation effects on UCP1 expression in Zucker rat MSCs

MSCs of inguinal WAT from *in vivo* melatonin-supplemented Zucker rats were isolated to study the effect of melatonin in the commitment of MSCs to the brown-like thermogenic adipocyte lineage. As shown in Fig. 3, a significantly increased UCP1 expression was measured by western blot after *in vivo* melatonin supplementation in MSCs from both lean (C-ZL, 1.241 ± 0.035 ; M-ZL, 1.592 ± 0.102 ; $P < 0.05$) and diabetic fatty rats (C-ZDF, 0.761 ± 0.022 ; M-ZDF, 1.230 ± 0.154 ; $P < 0.01$) compared to their respective controls, with no differences in UCP1 expression between MSCs from obese melatonin-supplemented rats and lean control ones. In addition, MSCs from unsupplemented obese rats showed less UCP1 expression levels than samples from unsupplemented lean rats (C-ZL, 1.241 ± 0.035 ; C-ZDF, 0.761 ± 0.022 ; $P < 0.01$; Fig. 3) showing the effect of obesity in the UCP1 expression of MSCs.

3.3 Characterization of MSCs isolated from inguinal fat of Zucker rats and human lipoaspirates

MSCs were isolated from the inguinal fat pad of Zucker rats and human lipoaspirates. Flow cytometry was carried out to

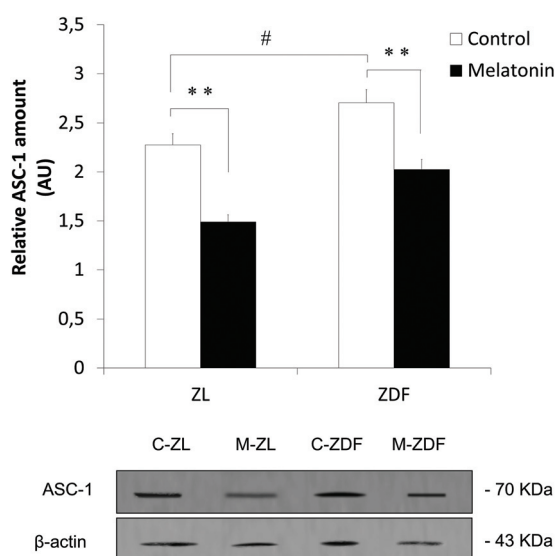


Fig. 2 Effects of *in vivo* melatonin supplementation on specific white adipocyte marker ASC-1 expression in inguinal adipose tissue from Zucker rats. ZL, Zucker lean rats; ZDF, Zucker diabetic fatty rats; C, control unsupplemented; M, melatonin-supplemented. All values are expressed as mean \pm SEM of three independent experiments in duplicate. ** $P < 0.01$; # $P < 0.05$.

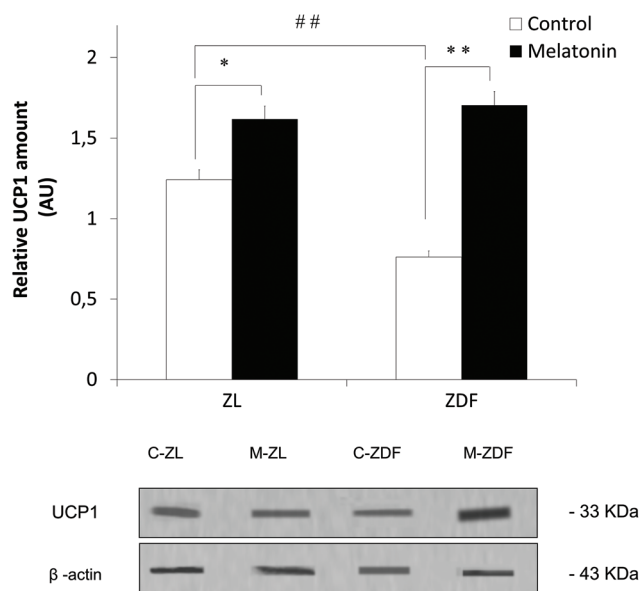


Fig. 3 Effects of *in vivo* melatonin supplementation on thermogenic protein UCP1 expression in MSCs of inguinal adipose tissue from Zucker rats. ZL, Zucker lean rats; ZDF, Zucker diabetic fatty rats; C, unsupplemented control; M, melatonin-supplemented. All values are expressed as mean \pm SEM of three independent experiments in duplicate. ** $P < 0.01$; * $P < 0.05$; ## $P < 0.01$.

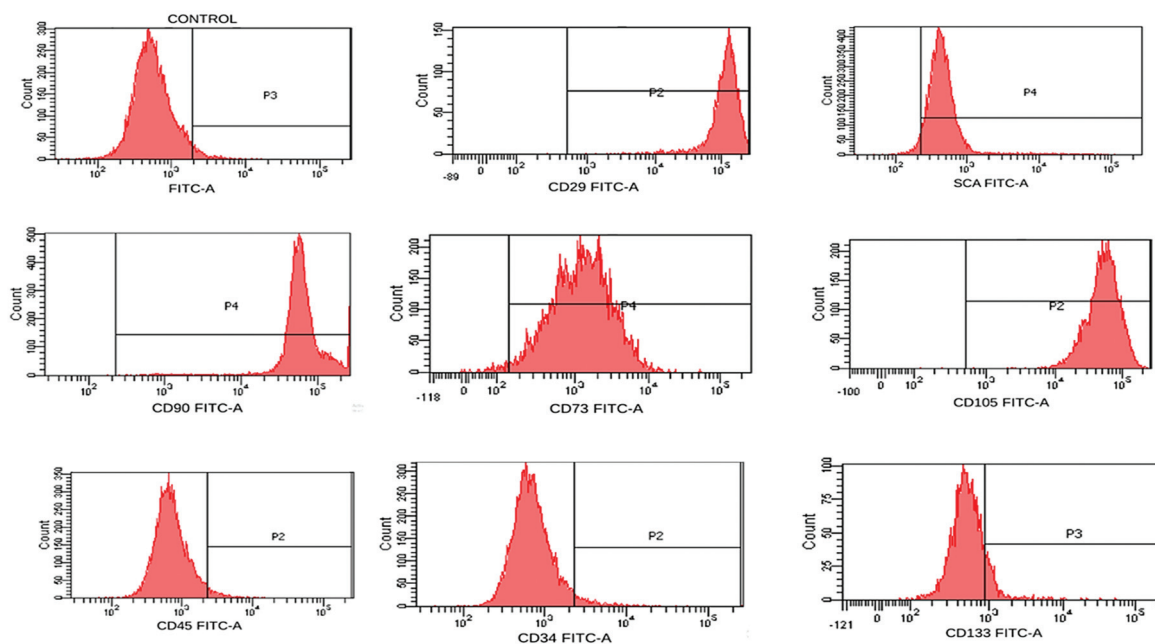


characterize the isolated cells using rat MSC specific markers CD-133 and stem cell antigen 1 (SCA-1), and CD-90, CD-73, CD-105, CD-34, CD-29, and CD-45 markers in both human and rat MSC. Cells were positive to CD-29, SCA-1, CD-105, CD-90 and CD-73, and negative to CD-45, CD-34 and CD-133 (Fig. 4A and B). Multipotency of isolated cells was tested by adipogenic, chondrogenic and osteogenic differentiation, and detected by oil red-O, toluidine, and alizarin red staining, respectively, after differentiation (Fig. 5A and B).

3.4 Analysis of differentiated adipocytes

After cell isolation from the inguinal WAT of ZDF rats and from human lipoaspirates, MSCs were subjected to adipogenesis assay to differentiate them into adipocytes (see Material and methods). Adipogenesis was successfully induced, and adipocytes were fully differentiated. Lipid droplet formation was examined by fluorescence microscopy (Fig. 6). AdipoRed staining enables the quantification of lipid droplets by measuring the absorbance at 520 nm, demonstrating a higher

A) ZDF rat



B) Human

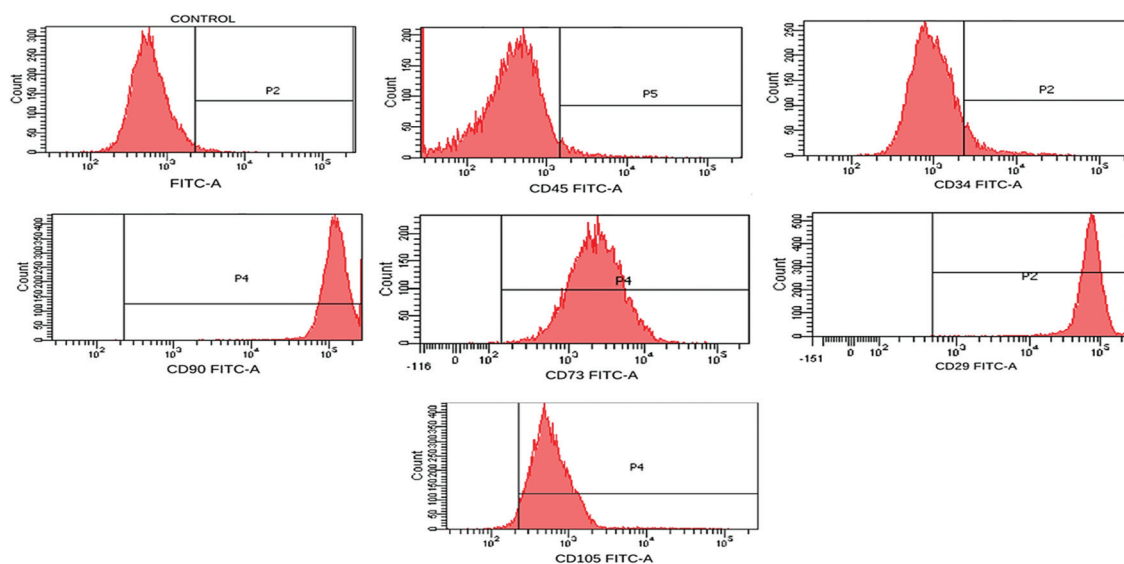
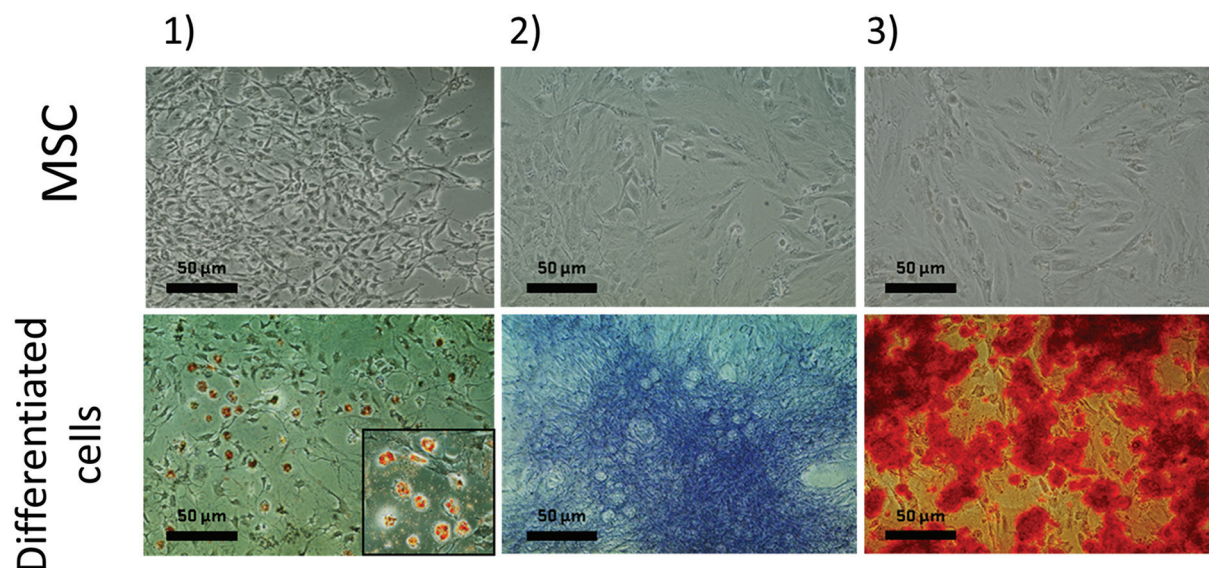


Fig. 4 Flow cytometry analysis of MSC isolated from (A) inguinal adipose tissue of Zucker rats and (B) human lipoaspirates. Samples without markers were used as a control. CD45, SCA-1, CD133, CD29 CD34, CD73, CD90 and CD105 were used as markers for MSC characterization.



A) ZDF rat



B) Human

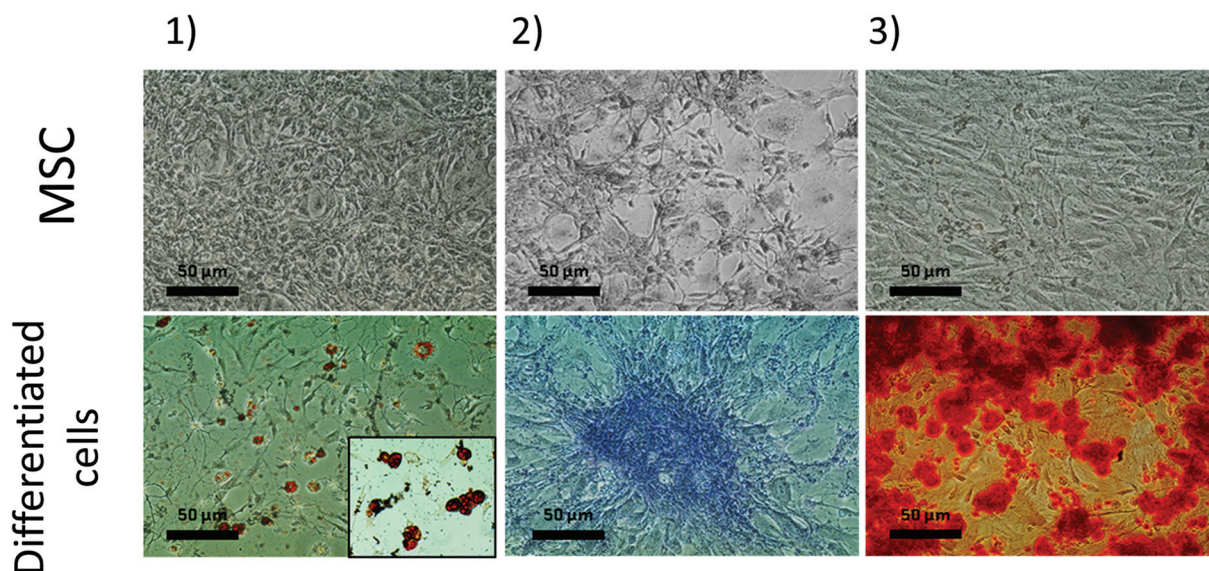


Fig. 5 *In vitro* characterization of cultured (A) Zucker rats and (B) human MSC. In both, in the upper panels, the images show MSC and in the lower panels, the differentiated cells from MSC after 21 days: (1) adipogenic, (2) chondrogenic and (3) osteogenic differentiation assay from optical microscopy. Magnification 40x. Scale bar 50 µm.

lipid content in differentiated adipocytes from both Zucker (MSC, 0.154 ± 0.015 ; differentiated adipocytes, 0.822 ± 0.029 ; $P < 0.01$; Fig. 6A) and human (MSC, 0.170 ± 0.027 ; differentiated adipocytes, 0.733 ± 0.051 ; $P < 0.01$; Fig. 6B) isolated cells.

3.5 *In vitro* melatonin treatment effects on UCP1, CITED1 and PGC1- α expression in ZDF rat MSCs

The thermogenic protein UCP1 expression was measured by western blot in ZDF MSC, untreated adipocyte (adipocytes)

and melatonin-treated adipocyte (Adipocytes + MLT) samples from both WMSC and bMSC iWAT from ZDF rat. No differences between WMSC and bMSC groups in UCP1 expression were found. Therefore, the WMSC and bMSC data were used interchangeably by recalculating the UCP1 expression of each group as the mean of the WMSC and bMSC expression. Melatonin *in vitro* treatment significantly induces UCP1 expression in differentiated adipocytes from ZDF MSCs (Adipocytes, 0.876 ± 0.044 ; Adipocytes + MLT, 1.063 ± 0.018 ; P



A) ZDF rat

B) Human

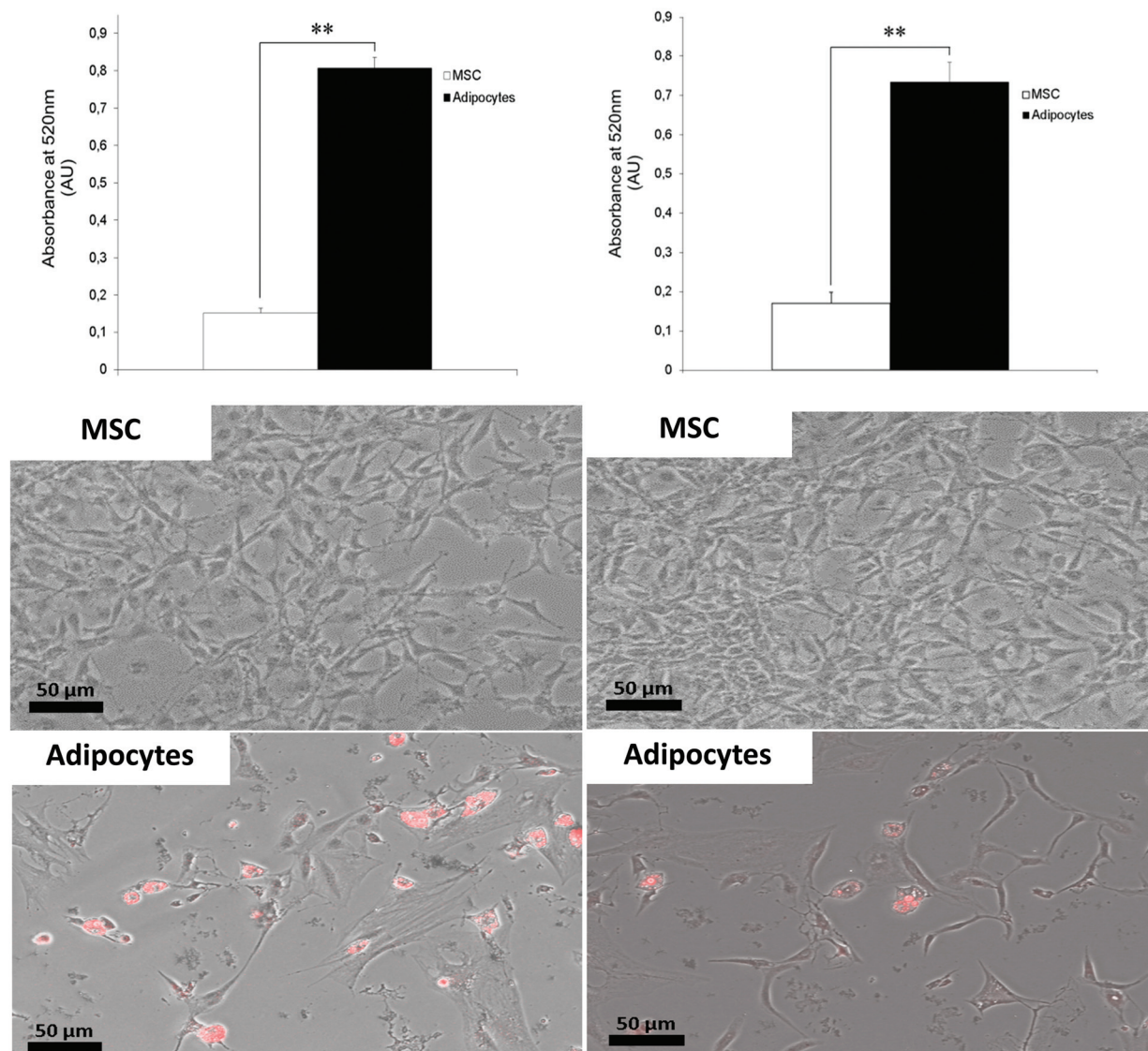


Fig. 6 Adipogenic differentiation of cultured MSC into adipocytes. Fluorescence microscopy images showed the lipid content in MSCs and differentiated adipocytes after *in vitro* melatonin treatment, measured by quantification of AdipoRed staining (absorbance at 520 nm) from (A) Zucker rats and (B) humans. All values are expressed as mean \pm SEM of three independent experiments in duplicate. MSC, mesenchymal stem cells; Adipocytes, differentiated adipocytes after 21 days. Magnification 63 \times . Scale bar 50 μ m. ****** P < 0.01.

< 0.05; Fig. 7) resulting in higher brown-like adipogenesis. No differences in UCP1 expression between MSCs and untreated differentiated adipocytes were found.

PGC1- α and CITED1 expression in ZDF MSCs, untreated adipocytes and melatonin-treated adipocytes was also evaluated by the western blot technique. WMSC and bMSC inguinal adipose tissue from ZDF rats were isolated to study the extent to which melatonin achieves commitment of MSCs to the beige phenotype adipogenic lineage. MSCs from bMSC present both higher PGC1- α (bMSC, 0.534 ± 0.026 ; WMSC, $0.476 \pm$

0.015 ; P < 0.05; Fig. 8A) and CITED1 (bMSC, 0.323 ± 0.009 ; WMSC, 0.236 ± 0.012 ; P < 0.05; Fig. 8B) expression levels than WMSC, suggesting a greater MSCs commitment to thermogenic beige adipocytes. After adipogenesis, no significant differences in CITED1 and PGC1- α expression between adipocytes differentiated from bMSC and WMSC were found regardless of the adipogenic differentiation assay. However, comparing groups, the melatonin-treated adipogenesis assay induces a significantly greater PGC1- α expression in differentiated ZDF adipocytes from both bMSCs (Adipocytes + MLT, 0.731 ± 0.043 ;



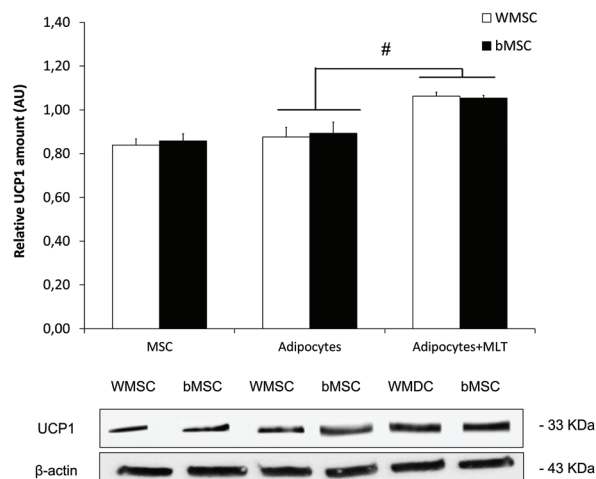


Fig. 7 Effects of *in vitro* melatonin treatment on thermogenic protein UCP1 in differentiated adipocytes from WMSCs and bMSC inguinal adipose tissue from ZDF rats. MSC, mesenchymal stem cells; Adipocytes, differentiated adipocytes after 21 days, Adipocytes + MLT, differentiated adipocytes after 21 days incubated with melatonin. All values are expressed as mean \pm SEM of three independent experiments in duplicate. # $P < 0.05$.

Adipocytes, 0.503 ± 0.034 ; $P < 0.05$) and WMSCs (Adipocytes + MLT, 0.724 ± 0.066 ; Adipocytes, 0.558 ± 0.054 ; $P < 0.05$), suggesting that melatonin increases the commitment of MSCs to differentiated brown-like adipocytes (Fig. 8A). Furthermore, the adipogenesis assay increases CITED1 expression in ZDF differentiated adipocytes compared to MSCs regardless of their origin, with significantly higher CITED1 expression in melatonin-treated adipocytes than in treated ones (MSC, 0.280 ± 0.052 ; Adipocytes, 0.528 ± 0.054 ; Adipocytes + MLT, 0.679 ± 0.038 ; on average; $P < 0.01$; Fig. 8B).

3.6 *In vitro* melatonin treatment effects on UCP1, CITED1 and PGC1- α expression in human MSCs

Western blot techniques were accomplished to measure UCP1, PGC1- α and CITED1 expression levels in lipoaspirate-derived human MSCs (hMSC), untreated hMSC-differentiated adipocytes (Adipocytes) and melatonin-treated hMSC-differentiated adipocytes (Adipocytes + MLT). *In vitro* melatonin treatment induces the expression of both thermogenic markers UCP1 (Adipocytes, 1.135 ± 0.038 ; Adipocytes + MLT, 1.284 ± 0.030 ; $P < 0.05$; Fig. 9A) and PGC1- α (Adipocytes, 3.130 ± 0.038 ; Adipocytes + MLT, 3.483 ± 0.130 ; $P < 0.05$; Fig. 9B) in hMSCs from healthy donors' lipoaspirates with significant expression differences between untreated and melatonin-treated differentiated adipocytes.

Moreover, the expression of the beige adipocyte marker CITED1 experiences a highly significant increase with melatonin treatment (Adipocytes, 0.645 ± 0.089 ; Adipocytes + MLT, 0.946 ± 0.078 ; $P < 0.01$; Fig. 9C). No significant differences between hMSCs and untreated adipocytes were found in any expression of the measured thermogenic proteins.

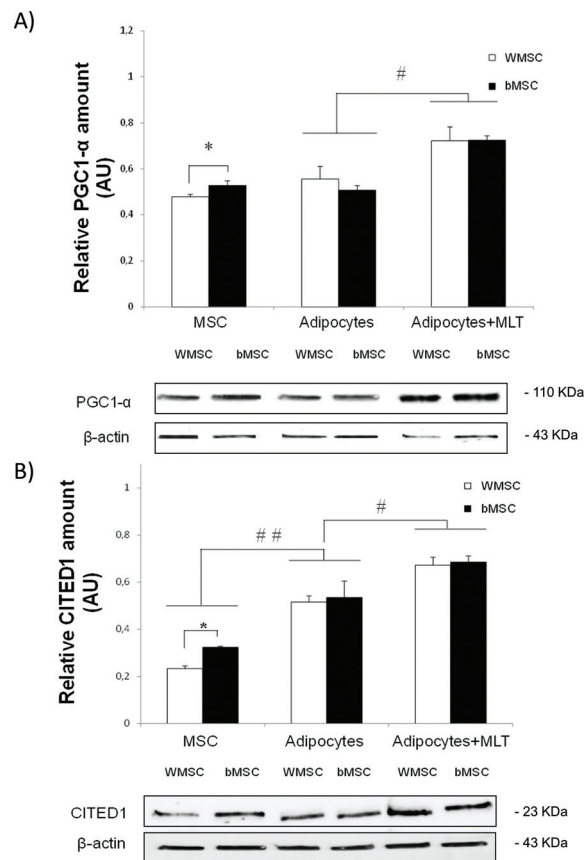


Fig. 8 Effects of *in vitro* melatonin treatment on brown-like cell marker (A) PGC1- α and (B) CITED1 expression in differentiated adipocytes from WMSCs and bMSC inguinal adipose tissue from ZDF rats. MSC, mesenchymal stem cells; Adipocytes, differentiated adipocytes after 21 days, Adipocytes + MLT, differentiated adipocytes after 21 days incubated with melatonin. All values are expressed as mean \pm SEM of three independent experiments in duplicate. ** $P < 0.01$; * $P < 0.05$.

4. Discussion

We previously reported that melatonin induces inguinal WAT browning in obese and diabetic Zucker rats, validating these results through the increased expression of UCP1. These results demonstrated that chronic oral melatonin drives WAT to brown multilocular fat function in ZDF rats.²⁵ In this study, our group also demonstrated through light microscopy morphological studies that unsupplemented ZDF rats present a smaller but larger number of adipocytes with a single fat drop and peripheral nucleus (white adipocytes) in the inguinal WAT of ZDF rats compared to the thin ones.²⁵ All these data are consistent with the lower fluorescence signal in unsupplemented ZDF rats and the partially recovered one in melatonin-supplemented ZDF rats by the appearance of multilocular cells (beiging) observed in this study. The increased expression of UCP1 is related to thermogenically active brown and beige adipose tissues.⁴⁷ However, beige/brite cells have a different gene expression pattern than white or brown fat cells.⁶ Indeed, CITED1 is known to be selectively expressed in beige cells and



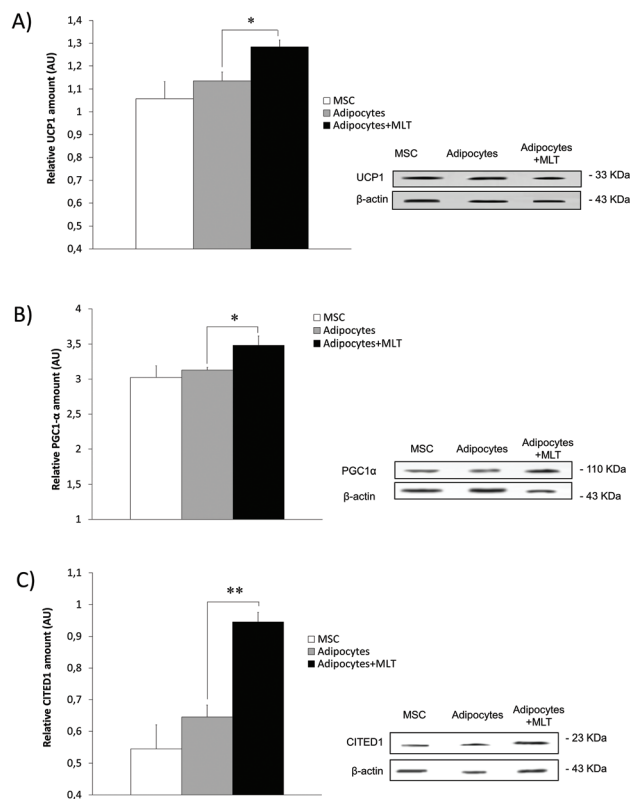


Fig. 9 Effects of *in vitro* melatonin treatment on thermogenic protein (A) UCP1, (B) PGC1- α and (C) brown-like cell marker CITED1 expression in differentiated adipocytes from hMSCs isolated from lipoaspirates' adipose tissue. MSC, human mesenchymal stem cells; Adipocytes, differentiated adipocytes after 21 days; Adipocytes + MLT, differentiated adipocytes after 21 days incubated with melatonin. All values are expressed as mean \pm SEM of three independent experiments in duplicate. ** $P < 0.01$; * $P < 0.05$.

to act as a transcriptional coactivator for estrogen receptors that were originally identified in a murine melanoma cell line.^{10,11} In contrast, ASC-1 is found to be a large surface marker for WAT due to its high expression in this tissue, with negligible or no expression in BAT.^{7,8}

Herein, we report for the first time that oral melatonin administration not only induces UCP1 and CITED1 positive beige adipocyte biogenesis in inguinal WAT from obese Zucker rats, but also decreases the ASC-1 expression in inguinal fat from both obese and lean Zucker rats, which confirm a reduced white adipocyte amount in inguinal WAT depots, in line with I. Murano *et al.* observations about the decline in the number of white adipose cells.⁴⁸ These results could explain the origin of melatonin-induced beige adipocytes by the partial transdifferentiation of white adipocytes to beige ones. This direct conversion process has been also demonstrated after cold acclimatization and concluded to be mainly mediated by the β -3 adrenergic receptor.³¹ However, more studies are needed to understand this biological process. Moreover, western blot analysis revealed that ASC-1 expression is higher in obese rats than in lean ones, indicating increased

inguinal WAT mass, typically observed in obese rodents.⁴⁹ Partial transdifferentiation shown alone cannot explain the elevated beige process observed in the inguinal WAT from ZDF rats after melatonin supplementation, suggesting a MSC *de novo* differentiation into beige adipocytes.

MSCs, which are multipotent stem cells, have the ability to turn into mature adipocytes in the final phase of differentiation under certain types of stimuli.⁵⁰ Therefore, as partial transdifferentiation was suggested in rats by the immunofluorescence assay, *de novo* differentiation studies of MSCs to mature adipocytes were performed to assess both hypotheses about the origin of melatonin-induced beige adipocytes in inguinal WAT of obese rats and in humans.

The isolation and characterization of inguinal fat and liposuction cells were performed by flow cytometry using, as proposed by the International Society for Cell Therapy (ISCT), positive MSC-specific surface markers such as CD-90, CD-73 and CD-105, and negatives like CD-34 and CD-45. In addition, other commonly measured surface markers were used in MSC characterization such as CD-29, a positive marker for both human and rat MSCs, SCA-1, a positive marker only for murine MSCs, and CD-133, a marker negative for MSC and only positive for hematopoietic stem cells.⁵¹ After characterization and the adipogenesis process, cells underwent in an *in vitro* melatonin treatment that mimicked the light-dark life cycle of circulating melatonin.^{52,53}

The high expression of UCP1 has been associated with thermogenic adipose tissue, as well as with brown adipocytes.^{6,37,41,47} The specific differentiation of MSCs to beige/brite adipocytes with rosiglitazone increases the expression of thermogenic genes such as UCP1 in rodents⁵⁴ and humans.^{55,56} The cold-induced browning of WAT deposits in rodents and humans is assessed by measuring the expression of UCP1.^{57,58} Cold exposure also induces MSC browning by increasing UCP1 expression in adipocytes differentiated from MSCs.⁵⁹ In this study, we report that *in vitro* melatonin treatment increases UCP1 expression in MSC-differentiated adipocytes, suggesting increased brown-type adipogenesis in human and rat-treated MSCs. Another study that compared the white and brown human adipogenesis of MSCs and umbilical cord stem cells showed no differences in the expression of UCP1 between MSCs and differentiated white adipocytes,⁶⁰ agreeing with the data observed in MSCs and differentiated adipocytes from inguinal WAT of ZDF rats and human lipoaspirates. Various studies in humans⁶¹ and rodents⁵⁹ revealed that MSCs and preadipocytes involved in adipogenesis showed the increased expression of thermogenic genes such as UCP1, indicating their thermogenic potential. These data are consistent with those obtained after *in vivo* melatonin supplementation reported in this research, in which MSC UCP1 expression increases in both lean and obese supplemented Zucker rats, showing a greater thermogenic potential for beige adipogenesis. Furthermore, we observed that the expression of UCP1 in unsupplemented obese rats was lower than that in unsupplemented lean rats, suggesting a decrease in the compromise in MSCs derived from ZDF rats and a lower



thermogenic activity of future differentiated adipocytes. In humans, subcutaneous WAT-derived MSCs from obese patients also exhibit less expression of PGC1- α , CITED1, and UCP1 than lean MSCs, with decreased changes in thermogenic gene expression between pre- and post-differentiation.⁶² In this study, we also report for the first time that melatonin induces the expression of thermogenic proteins UCP1, CITED1 and PGC1- α in hMSC-derived adipocytes, revealing the role of melatonin in human *de novo* beige adipogenesis. Although some studies demonstrate the key role that the Wnt signaling pathway plays in the melatonin-induced differentiation of MSCs through the modulation of transcriptional activation of peroxisome proliferator-activated receptor γ (PPAR γ),⁶³ SIRT1 being the key nutrient and energy sensor underlying Wnt/ β -catenin pathway activation. A recent study also showed that PDGFR α positive progenitors are crucial for *de novo* brown adipogenesis in interscapular adipose tissue under cold exposure, being also a good candidate target for adipogenesis regulation in the development of adipose tissue.⁴⁴ Further studies are needed to better understand the mechanism by which melatonin could have a decisive effect on MSC involvement with thermogenically active beige adipocytes *in vivo* increasing MSC commitment.

To evaluate the beige biogenesis potential of melatonin in cultured MSCs, we performed the characterization of differentiated adipocytes by measuring CITED1 as a specific beige cell marker.⁹ Moreover, we determined the PGC1- α expression commonly associated with adaptive thermogenesis and mitochondrial biogenesis in rodents^{64,65} and humans,⁶⁶ besides being highly expressed in brown-like adipocytes. CITED1 and PGC1- α expression was higher in melatonin-treated differentiated adipocytes compared to untreated ones and undifferentiated MSCs in both studied species. This reaffirms the hypothesis that melatonin increases specific beige adipogenesis. Other studies demonstrated that cold⁵⁹ and adrenergic stimulation by sirutin 1 (SIRT1)^{67,68} or bone morphogenetic protein 7 (BMP7)⁶⁹ also promote the thermogenic gene program, enhancing UCP1, PGC1- α and PR-domain containing 16 (PRDM16) expression, and high expression of beige adipocyte markers like CITED1, purinergic receptor P2X 5 (P2RX5) and CD137 in differentiated adipocytes. Recent studies in MSC differentiation to BAT in humans revealed that the increased expression of PGC1- α in MSCs is related to higher mitochondrial mass and activity, with higher UCP1 expression (mitochondrial biogenesis), typically observed in beige and brown adipocytes.⁷⁰ They also found by microarray analysis that PGC1- α expressing MSCs has a shorter average linkage distance to mature brown adipocytes, showing a greater commitment to brown-like adipogenesis.

CITED1 was highly expressed in adipocytes differentiated from untreated inguinal MSCs, and the data are consistent with many studies in mice where the authors demonstrated that subcutaneous and especially inguinal adipose tissue are more prone to become thermogenic beige adipose tissue.^{71,72} We also observed that the expression of CITED1 and PGC1- α was significantly higher in MSCs derived from beige deposits

of the inguinal fat pad than in white ones, which suggests a greater commitment of MSCs with the differentiation of thermogenic beige adipocytes in the bMSCs than WMSCs. This could be explained by the epigenomic memory found in beige adipocytes,⁴⁶ which could be present in BAT and explain similar data obtained in mice⁷³ and human studies^{60,74,75} in which derived MSCs from BAT were found to express higher levels of thermogenic proteins. A long-term cold exposure study in mice revealed that *de novo* beige biogenesis was favored from adipogenic precursor cells,⁷⁶ suggesting that the beige biogenesis mechanism, either by darkening of pre-existing mature adipocytes or *de novo* differentiation, also depends on the frequency and duration of stimuli such as cold.⁷⁷ Therefore, these are two key variables in the melatonin regulation of MSC adipogenesis as a promising therapeutic target for obesity control. In addition, other effects of melatonin were recently demonstrated on mesenchymal stem cells derived from the umbilical cord blood of obese rodents that reduce cholesterol accumulation and prevent apoptosis of MSCs, supporting the therapeutic role of melatonin against obesity.⁷⁸ A recent meta-analysis of the effects of melatonin in human obesity studies showed that, despite the evidence found in the energy balance of rodents and humans,^{79,80} the real impact of melatonin on body weight loss remains unclear due to the inconsistent melatonin treatment schedule regarding duration and low dose (0.01–0.1 mg kg⁻¹ day⁻¹) in all studies. Overall, these data provide evidence that, under melatonin treatment, the *de novo* differentiation of beige cells in ZDF rats and humans probably occurs in conjunction.

Conclusions

In conclusion, with these results, it is demonstrated that the administration of melatonin to obese and diabetic ZDF rats induces beige biogenesis by the partial transdifferentiation of white to beige adipocytes, reducing the amount of white adipose cells and the differentiation of MSCs in the inguinal adipose tissue. In addition, melatonin plays a key role in the beige adipogenesis of human MSCs, promoting MSC engagement with the brown-type thermogenic adipocyte lineage and enhancing the expression of thermogenic markers. Therefore, as a thermogenic inducer of browning, melatonin could be a safe and effective strategy to combat obesity and metabolic diseases. More studies on the nuclear-mitochondrial connection are needed to better understand the thermogenic pathways involved, with melatonin receptors and mitochondrial DNA being fundamental study targets to identify mitochondrial-nucleus communication in melatonin treatment for human obesity and diabetes.

Ethical statement

The study was conducted according to the guidelines of the Declaration of Helsinki and approved by the Ethical



Committee of the University of Granada (Granada, Spain) (4 09-2016-CEEA and 02/022010). Informed consent was obtained from all subjects involved in the study.

Author contributions

AA conception and design, financial support, manuscript writing, final approval of manuscript. DS, MNA and MC collection and assembly of data, data analysis and interpretation, manuscript writing. GFV, MJOL, AMH and AMC manuscript writing. All authors have read and agreed to the published version of the manuscript.

Conflicts of interest

The authors declare that there are no conflicts of interest.

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