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# Beta-hydroxybutyrate dampens adipose progenitors' profibrotic activation through canonical Tgf $\beta$ signaling and non-canonical ZFP36-dependent mechanisms

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## ABSTRACT

Adipose tissue contains progenitor cells that contribute to beneficial tissue expansion when needed by *de novo* adipocyte formation (classical white or beige fat cells with thermogenic potential). However, in chronic obesity, they can exhibit an activated pro-fibrotic, extracellular matrix (ECM)-depositing phenotype that highly aggravates obesity-related adipose tissue dysfunction.

**Objective:** Given that progenitors' fibrotic activation and fat cell browning appear to be antagonistic cell fates, we have examined the anti-fibrotic potential of pro-browning agents in an obesogenic condition.

**Results:** In obese mice fed a high fat diet, thermoneutral housing, which induces brown fat cell dormancy, increases the expression of ECM gene programs compared to conventionally raised animals, indicating aggravation of obesity-related tissue fibrosis at thermoneutrality. In a model of primary cultured murine adipose progenitors, we found that exposure to  $\beta$ -hydroxybutyrate selectively reduced Tgf $\beta$ -dependent profibrotic responses of ECM genes like *Ctgf*, *Loxl2* and *Fn1*. This effect is observed in both subcutaneous and visceral-derived adipose progenitors, as well as in 3T3-L1 fibroblasts. In 30 patients with obesity eligible for bariatric surgery, those with higher circulating  $\beta$ -hydroxybutyrate levels have lower subcutaneous adipose tissue fibrotic scores. Mechanistically,  $\beta$ -hydroxybutyrate limits Tgf $\beta$ -dependent collagen accumulation and reduces Smad2-3 protein expression and phosphorylation in visceral progenitors. Moreover,  $\beta$ -hydroxybutyrate induces the expression of the *ZFP36* gene, encoding a post-transcriptional regulator that promotes the degradation of mRNA by binding to AU-rich sites within 3'UTRs. Importantly, complete *ZFP36* deficiency in a mouse embryonic fibroblast line from null mice, or siRNA knock-down in primary progenitors, indicate that *ZFP36* is required for  $\beta$ -hydroxybutyrate anti-fibrotic effects.

**Conclusion:** These data unravel the potential of  $\beta$ -hydroxybutyrate to limit adipose tissue matrix deposition, a finding that might be exploited in an obesogenic context.

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**Keywords** Adipocyte; Progenitors; Fibrosis; Extracellular matrix

## 1. INTRODUCTION

Increasing fat mass over time favours progression to metabolic diseases and type-II diabetes. Maladaptive metabolism in obesity relies on chronic low-grade inflammation and subsequent disruption of insulin sensitivity [1]. It is linked to metabolic inflexibility of the expanded fat pads, which acquire features of a stiff, extracellular matrix-embedded fibrotic tissue, unable to sustain efficient lipid sequestration [2]. Adipose tissue (AT) fibrosis has strong deleterious impacts on glucose homeostasis, demonstrated in obese subjects [3,4] and in mouse models [5,6]. Conversely, thermogenic AT (brown or beige) is highly efficient in glucose uptake and lipid utilization [7,8], therefore a

beneficial actor in blood glucose control [9,10]. Pathways controlling acquisition of a thermogenic fat cell phenotype are linked to the induction of a specific mitochondrial program dependent on PPAR $\gamma$ -associated transcriptional regulators like PRDM16 and PGC1a [11]. The recruitment of a pool of thermogenic adipocytes to target obesity-associated metabolic impairment has now become a therapeutic strategy under active research.

The source of brown-like and white adipocytes is a bulk of progenitors residing within the AT stroma-vascular fraction. The global adipose progenitor population has different propensities for beige/white adipogenesis, and can also develop into a pro-fibrotic fibroblastic phenotype, underlining a dual fibro/adipogenic potential highly

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dependent on the tissue microenvironment. Although highly heterogeneous and yet incompletely characterized, the pool of AT progenitors (Cd31-, Cd45-) shares PDGFR $\alpha$  expression as a common cell surface marker [12]. Using this marker for functional progenitor targeting, we observed that fibrogenic responses to high fat diet, and fat browning potential of mouse subcutaneous adipose tissue, were inversely related pathways [6]. In line, another report demonstrated an inverse relationship between mouse fat browning and AT fibrosis, mediated by a PRDM16-dependent signal released from the adipocytes [13]. Thus, progenitors' fibrotic transformation and brown-like adipogenesis may involve antagonistic cell fates, associated with opposite consequences on whole body metabolic regulation.

A better understanding of how adipose progenitors orient cell choices towards fibrogenesis is essential for the development of new approaches to limit metabolic deterioration in obesity. A switch in the abundance of CD9-expressing visceral progenitors has been reported to promote pro-fibrotic activation [4]. It is also likely that the obese AT pro-inflammatory microenvironment triggers fibrogenesis at the expense of adipogenesis, linked to disruption in Tgf $\beta$  (Transforming growth factor) family members balance [14]. Moreover, the high demand for lipid storage in obesity might lead to preadipocytes premature exhaustion and senescence, precluding an adequate supply of new adipocytes [15].

In this study, we have explored the paradigm of pro-fibrotic versus adipogenic development of progenitors in the context of the obese AT. In particular, we demonstrate aggravated fibrosis in obese high fat diet-fed mice in which brown/beige fat dormancy is shaped by raising at thermoneutrality. We also demonstrate the anti-fibrotic properties of the fat browning factor  $\beta$ -hydroxybutyrate in primary Tgf $\beta$ -induced cultured progenitors from subcutaneous or visceral tissues. In addition to blunting of Smad/Tgf $\beta$  signalling by  $\beta$ -hydroxybutyrate in visceral progenitors, we unravel a novel mechanism linked to ZFP36/Tristetraprolin, a post-transcriptional regulator of mRNA stability, which is induced by  $\beta$ -hydroxybutyrate, and restrains fibrotic activation of subcutaneous progenitors.

## 2. MATERIALS AND METHODS

### 2.1. Materials

Collagenase A (R11088793001) was purchased from Roche, DMEM/Glutamax (61,965-026) from Gibco and foetal calf serum (CVFSVF00-01) from Eurobio Scientific. Sodium 3- $\beta$ -Hydroxybutyrate (54,965) and L-(+)-lactic acid (L1750) were obtained from Sigma. Tgf $\beta$ 1 (human recombinant, ref 130-126-724) was from Myltenyi Biotech, Bmp4 (Human recombinant, ref PHC9534) from Invitrogen.

### 2.2. Patient studies

Human samples used were obtained from patients with obesity (N = 30, age 42.5 $\pm$ 14.5, 10 men, 20 women, body mass index 44.0 $\pm$ 6.5 kg/m<sup>2</sup>) involved in the bariatric surgery program at the Nutrition Department of Pitié-Salpêtrière hospital, France. Ten patients (7 women, 3 men) had normal glucose control, while the 20 others were either diabetic or glucose intolerant. Among diabetics, 4 were treated by GLP1 analogs, 2 by insulin, 5 by metformin. Six patients (3 men, 3 women) received statins to treat hyperlipemia. Eleven (7 women, 4 men) were on anti-hypertensive drugs. Blood samples were collected during preoperative hospital examination after an overnight fast. All patients received nutritional information and are encouraged to healthy diets. No specific diet pattern is recommended before surgery. Patients were part of a follow-up cohort with tissue bio banking during the surgery and benefited fibrosis scoring of subcutaneous AT based

on Picrosirius Red labelling of paraffin-embedded samples as described previously [16]. Briefly, label detection thresholds were adjusted with an image-analysis module using Calopix software (TRIBVN) and manual delimitation of red-stained AT, avoiding blood vessels and staining artifacts was performed. Label quantification was expressed as the ratio of red-stained to total tissue area. A combined semi-quantitative SCAT fibrosis score (FAT score) was then attributed to each patient according to the following rules: stage 0: no perilobular staining (thickness less than the diameter of average adipocyte) and no pericellular fibrosis (no accumulation of collagen around adipocytes localized in the depth of the AT lobules). Stage 3: severe parabolular fibrosis (thickness similar to the diameter of two average adipocytes) and severe paracellular fibrosis (thick labeling around adipocytes and presence of within label-trapped adipocytes). Ethical approval was obtained from the Research Ethics Committee of Hôtel-Dieu Hospital (CPP Ile-de-France N° 1). Informed written consent was obtained from all subjects and the protocol was registered on <http://www.clinicaltrials.gov> (NCT01655017, NCT00476658).  $\beta$ -Hydroxybutyrate was measured on serum aliquots (5  $\mu$ l) stored at -80C with a colorimetric assay kit (Abnova KA0854).

### 2.3. Mouse studies

Animal studies were conducted in agreement with ARRIVE guidelines, approved by local ethical committee (n°31,719). 4–6 week old C57Bl6/J mice were fed a high fat diet (Research Diet, D12492i) with free access to drinking water. They were housed (maximum of 5 mice per cage) at room temperature (22 °C) or at thermoneutrality (30 °C) in a climate chamber (TSE, PhenoMaster), with matched day–night light cycles and controlled humidity. Body composition was analyzed by nuclear magnetic resonance (Bruker Mouse Minispec, LF90). AT from visceral (gonadal), subcutaneous (inguinal) and interscapular brown fat (BAT) were frozen in liquid nitrogen or incubated with collagenase as described [6].

### 2.4. Primary cell culture

Six female mice fed on a standard chow diet were used for each cell preparation. Minced AT were incubated in collagenase dissociation medium for 30 min (visceral AT) or 1 h (Subcutaneous AT) under lateral shaking (100 rpm) at 37 °C. After filtration through a 100  $\mu$ m cell strainer, medium was adjusted to 30 ml with PBS, cells were pelleted (10 min at 1500rpm), and suspended into 12 ml of DMEM/10% fetal calf serum. 0.5 ml was plated into 12-well culture plates in a 95%–5% air-CO<sub>2</sub> humidified atmosphere at 37 °C. After 4 h, non-attached cells were eliminated by 3 washes with PBS, and refed in fresh medium. The next day, all cultures were more than 80% confluent, and  $\beta$ -hydroxybutyrate ( $\beta$ OHB) (or other browning agents) was added alone 24 h prior to TGF $\beta$  was introduced and maintained for additional 3 days. This combinatory treatment was defined to mimic the transition from a healthy pro-browning to a pro-fibrotic AT microenvironment.

### 2.5. Transfection with siRNA

Cells in DMEM 10% fetal calf serum were incubated with 20 nM siRNA duplexes (SR426376) or control scrambled sequences (SR0003) from Origene, using the lipofectamine RNAiMax transfection reagent (ref 13,778- Invitrogen). Cell treatment with effectors was performed on the next day in fresh medium.

### 2.6. A ZFP36-deficient mouse fibroblast

Cell line and its matched WT control line were originally derived from ZFP36 -/- and +/+ embryos [17], and were cultured in DMEM high

1 glucose medium containing 1% penicillin/streptomycin and 10% fetal  
2 calf serum.

### 3 2.7. Gene and protein expression

4 Total RNA was extracted using RNeasy Mini kit (Qiagen, 74,104), and  
5 500 µg was used for reverse transcription, followed by Quantitative  
6 real-time PCR using SyBr Green Master Mix (Applied Biosystems). All  
7 primer pairs were validated with cDNA calibration curves and a unique  
8 amplification product. For protein analysis, lysates in RIPA buffer  
9 (Sigma Aldrich, R0278) containing proteinase inhibitors (Complete  
10 Mini, Roche) were prepared (Bertin Technologies, Precellys 24) and  
11 cleared for 10 min at 10,000 rpm. Protein concentration was  
12 measured (ThermoFisher Scientific, BCA assay). Western blotting was  
13 performed as described [18].

### 14 2.8. Bioinformatics

15 We downloaded six data sets from publicly available reports [19–21]  
16 detailing information from chow-fed mice of different sexes, devel-  
17 opmental states or adipose tissue anatomical locations. To keep a  
18 global overview of all datasets, we did not remove any genes from any  
19 of the six datasets, and kept all mitochondrial genes. In order to  
20 combine several scRNAseq experiments from different mouse studies  
21 while minimizing batch effects, we used Batch balanced K nearest  
22 Neighbours method (BBKNN) [22] available in the Scanpy Package to  
23 convert the distance between members of the same cluster into  
24 projectable connectivity. We then performed a Leiden Clustering  
25 (<https://www.nature.com/articles/s41598-019-41695-z>),  
26 which defined groups with similarity relationships. For each cluster, a ranking  
27 of genes defining the cluster was established. Based on a list of genes,  
28 we looked at the normalized expression of these genes in each of the  
29 clusters and performed stacked violin plots.

### 30 2.9. Statistical analysis

31 Cell culture experiments were performed in triplicate wells, repeated  
32 at least three times with independent cell preparations. Mean values from  
33 independent experiments were compared by Student's *t* test with a  
34 significance threshold at  $p < 0.05$ . In mouse experiments, differences  
35 in mean values from individual mice (6–10 per group) were evaluated  
36 by Student's *t* test. Spearman correlations were considered significant  
37 when  $p < 0.05$ .

## 38 3. RESULTS

### 39 3.1. Thermogenic fat dormancy by thermoneutral housing 40 aggravates obesity-dependent extracellular matrix (ECM) gene 41 expression in mouse AT

42 Cold exposure that activates beige adipocyte biogenesis reduces ad-  
43 ipose tissue fibrosis and improves systemic glucose homeostasis in  
44 mice [5,23]. To examine if shutting down of thermogenic activity could  
45 reciprocally increase the severity of high fat diet (HFD)-induced AT  
46 fibrosis we compared HFD mice (both sexes) housed at either ther-  
47 moneutrality (TN, 30 °C) or room temperature (RT, 20–22 °C). While  
48 pre- or post-HFD mean body weights did not significantly differ be-  
49 tween groups (Suppl Fig. 1A), TN female mice gained more weight  
50 than RT females, close to growth rates of RT males (Suppl Fig. 1B).  
51 Moreover, 9 weeks on HFD at TN instead of RT significantly increased  
52 fat mass in female mice (Suppl Fig. 1C). Thus, TN housing combined  
53 with HFD amplified fat accretion, more markedly in females than in  
54 males.

55 Compared to their conventionally raised counterparts fed HFD, TN mice  
56 expressed lower levels of BAT *Ucp1* mRNA (Figure. 1A), and developed

57 more inflamed BAT, with higher expression of *Ilf6*, *Tnfrα* and *Cd68*  
58 (Figure. 1B). Moreover, a number of genes involved in ECM develop-  
59 ment were overexpressed in TN compared to RT. Specifically, TN  
60 enhanced genes encoding fibrillary collagens (*Col1a1*, and *Col3a1*),  
61 ECM modifying enzymes such as *Loxl2* (a collagen/elastin crosslinker),  
62 and the matrix metalloproteinase regulator *Timp1* (Figure. 1C) which  
63 are implicated in the development of AT fibrosis [24]. Interestingly, BAT  
64 *Ucp1* negatively correlated with *Col1a1* and *Col3a1* expression (Figure.  
65 1D). Comparison of BAT histology showed bigger lipid droplets in TN  
66 mice, as expected (Figure. 1E). At low magnification, we noticed the  
67 presence of prominent white spans formed with α-cellular material  
68 indicative of deposited extracellular matrix delimiting lobules in TN  
69 BAT. Semi-quantitative analysis showed significantly increased num-  
70 ber of lobules in TN compared to RT samples (Figure. 1F). Thus, TN  
71 housing amplified the HFD-dependent ECM gene fibrotic program and  
72 matrix deposition in BAT.

73 In subcutaneous fat (SCAT), we found increased collagen1 protein  
74 content in TN versus RT mice (Figure. 1G). Although the topology of  
75 collagen accumulation in TN mice SCAT was not determined in the  
76 present study, we previously observed that HFD mice at RT prefer-  
77 entially accumulated collagen as perilobular fibrosis [6]. SCAT is a  
78 depot in which conventional white and brown-like fat cells are mixed,  
79 resulting in highly variable *Ucp1* mRNA expression among individual  
80 mice. A lower proportion of TN mice expressed high (above median)  
81 *Ucp1* mRNA, indicating a reduced propensity for SCAT beiging at TN  
82 (Figure. 1H). Among SCAT expressing *Ucp1* above the median, *Ucp1*  
83 mRNA negatively associated with *Col3a1* and *Fn1* (Figure. 1I). Thus,  
84 thermoneutral housing of mice showed an *Ucp1*-dependent inverse  
85 link to ECM gene patterns, which is consistent with antagonistic  
86 regulation of fibrotic and browning programs.

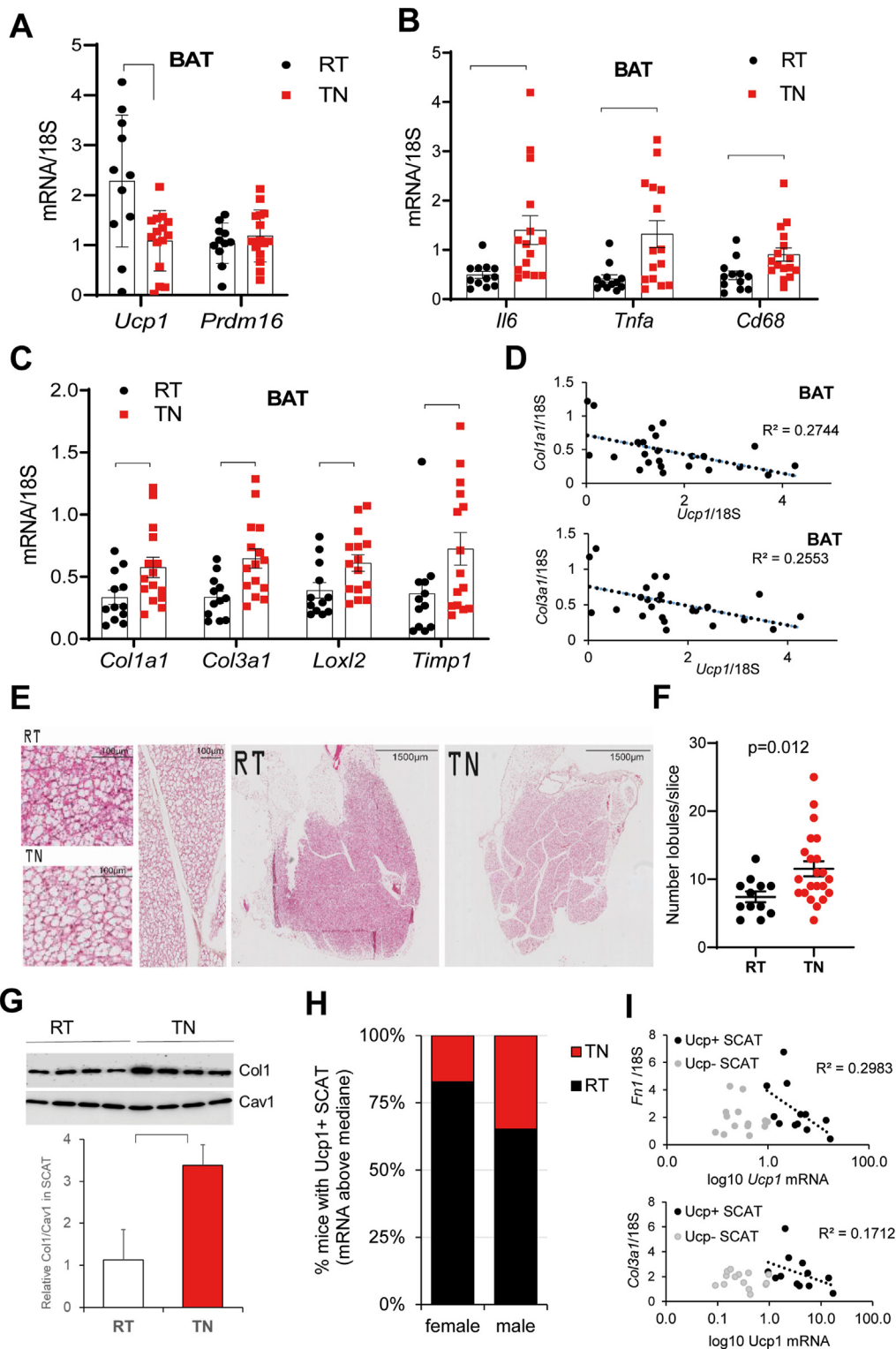
### 87 3.2. Reciprocal beiging and fibrotic responses of progenitors in a 88 simplified culture system

89 As AT resident progenitors are a common source of both beige fat cells  
90 or profibrotic fibroblasts, we aimed to establish a simplified *in vitro*  
91 system to investigate fibrotic versus beige fat development. We used  
92 the stroma-vascular cell fraction of mouse AT from subcutaneous  
93 (SCAT) or visceral (VAT) fat depots, known to exhibit common sus-  
94 ceptibility but distinct efficiencies for beiging or fibrotic development  
95 *in vivo*. Considering high heterogeneity of adipose progenitor cell  
96 population revealed by single cell sequencing [19–21], we aimed to  
97 minimize any drift in relative abundance of progenitor subsets that  
98 might occur during *in vitro* propagation. Cells recovered from colla-  
99 genase digestion were plated at high density (80% confluence after 4-  
100 hour attachment), which ensured minimal proliferation and limited  
101 composition shifts from potential differences in proliferative rates  
102 among progenitor subtypes. Moreover, only primary first-passage cells  
103 were used, and culture duration did not exceed 5 days.

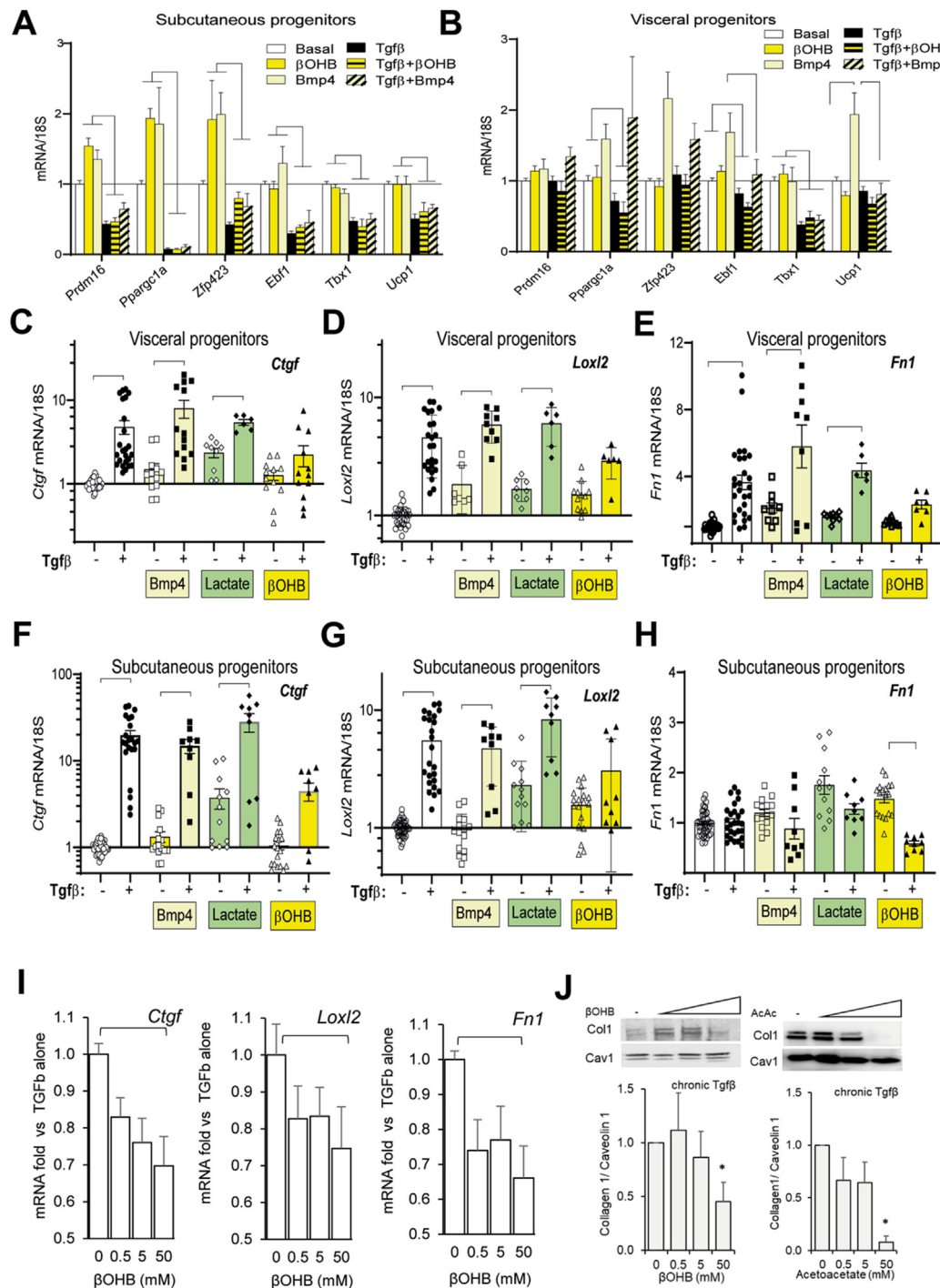
104 In this system, we tested cell responses to the canonical pro-fibrotic  
105 agent Tgfβ, as well as to previously characterized browning factors,  
106 i.e., β-hydroxybutyrate (βOHB) and Bone Morphogenic Protein 4  
107 (Bmp4) [13,25]. All progenitors responded to Tgfβ exposure by marked  
108 morphological changes, with elongated cell shapes and dense cell-  
109 cell contacts (Suppl Fig. 2A). As expected, pro-fibrotic Tgfβ treat-  
110 ment upregulated the expression of ECM marker genes (i.e. *Col1a1*,  
111 *Ctgf*, *Loxl2*) in progenitors derived from either visceral (Suppl Fig. 2B)  
112 or subcutaneous (Suppl Fig. 2C) fat. *Fn1* (encoding the abundant  
113 matrix protein fibronectin) and *Col3a1* mRNAs were up-regulated by  
114 Tgfβ in visceral progenitors only. We also observed that lactate  
115 addition in the culture medium stimulated ECM gene expression, with  
116 synergistic effect to Tgfβ.



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**Figure 1:** Thermoneutral housing aggravates HFD-induced adipose tissue ECM gene program in mice. Mice were housed for 9 weeks at room temperature (RT, 22 °C) or, thermoneutrality (TN, 30 °C) and received ad libitum HFD feeding. **A-D:** Brown adipose tissue (BAT) gene expression after 9 weeks on HFD was evaluated by Real time Q-PCR and normalized to 18 S. Values from individual mice are plotted, parentheses indicate significant differences between groups assessed by Student's t test. **E:** Hematoxylin/eosin BAT slices from RT and TN mice at different focus level (see scale bar in each image). Left panels show increased lipid droplet size in TN compared to RT (left panels). White spans consisting of acellular material (vertical image) were identified in whole tissue scans, delimiting lobules. **F:** Semi quantitative analysis of lobule number. Each point is from individual mice. **G:** Western blot analysis of SCAT Collagen1 content in RT and TN mice. Four tissues from individual mice are shown in each group; signals are normalized to Caveolin 1 for quantification. Parenthesis indicates significant difference ( $p < 0.05$ ) by t test. **H:** Proportion of mice with *Ucp1* positive SCAT (defined as above median *Ucp1* mRNA value) in RT and TN groups. P value = 0.035 by Chi2 Test. **I:** Correlation between *Ucp1* expression in SCAT and ECM genes (*Fnr1*: upper panel, *Col3a1*: lower panel). Significant correlation ( $p < 0.05$ ) is found in *Ucp1*+ SCAT (black) but not *Ucp1*- SCAT (grey). (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

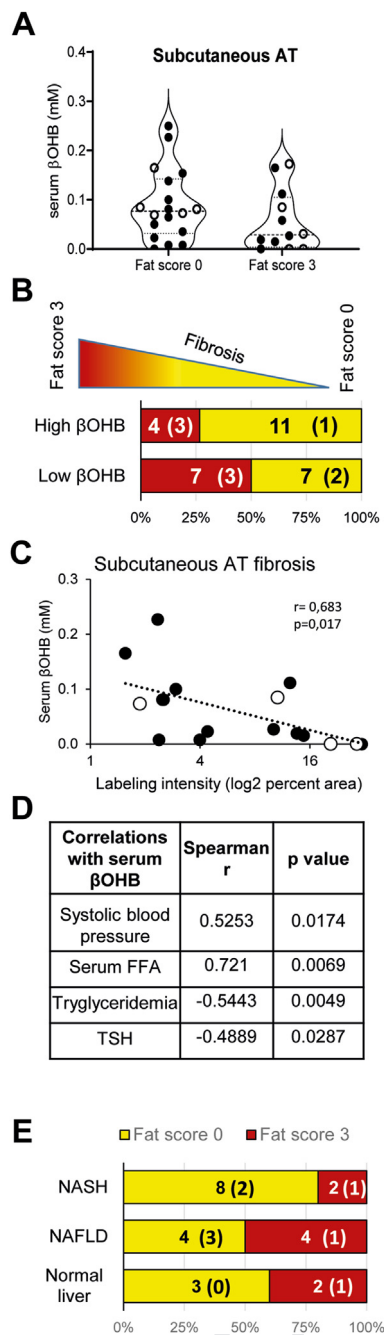


**Figure 2:**  $\beta$ OHB counteracts Tgfb-mediated stimulation of ECM genes. Combinatorial treatment with browning agents and Tgfb of visceral (A) or subcutaneous progenitors (B) on the expression of browning genes.  $\beta$ OHB (50 mM) or Bmp4 (10 ng/ml) was added 24 h after plating and Tgfb (10 ng/ml) was added the following day, until day 5. Bars are mean  $\pm$  sem of three independent cell preparations. The same experimental scheme was used in (C–H), to assess *Ctgf*, *Loxl2* and *Fn1* mRNA responses. Each point is an individual well from triplicates in at least 4 independent cell preparations. I: Dose-dependent response of *Ctgf*, *Loxl2* and *Fn1* gene expression to  $\beta$ OHB in the presence of Tgfb in primary subcutaneous progenitors. Parentheses indicate significant differences between conditions, by Student's t test. J: Western blot analysis of Collagen1 protein expression in Tgfb-stimulated visceral progenitors in the presence of  $\beta$ OHB or Acetoacetate.  $\beta$ OHB or Acetoacetate was added 1 day post plating and was maintained until cell harvest (Day 5). Chronic Tgfb stimulation started from Day 2 to Day 5. Collagen1 antibody was from Proteintech, ref14695-1. Representative blot is shown, with quantification from 2 to 4 independent cell preparations.

As expected, progenitor response to browning agents depended on fat tissue origin. Indeed, visceral progenitors (Suppl Fig. 2D) responded to Bmp4 by inducing *Ppargc1a* mRNA, a master regulator of the

mitochondrial program, as well as *Zeb1*, *Zfp423*, *Ebf1* and *Ucp1*.  $\beta$ OHB treatment had limited effects on visceral progenitors and only slightly increased *Zeb1* mRNA. Subcutaneous progenitors (Suppl Fig. 2E)

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**Figure 3:** Serum  $\beta$ OHB levels negatively associate with subcutaneous adipose tissue fibrosis in patients with obesity. **A:** Serum  $\beta$ OHB levels were assessed in a group of 29 obese patients eligible for bariatric surgery, in which the intensity of subcutaneous fat fibrosis had been scored as previously described. Men are represented as open symbols. **B:** Patient stratification according to serum  $\beta$ OHB (relative to median value) indicates preferential clustering of patients with low Fat score in the high  $\beta$ OHB group. Numbers represent the total number of patients in stratification groups, with the number of men in parenthesis. **C:** Negative association (Spearman correlation) of serum  $\beta$ OHB levels and intensity of ECM labelling by Picrosirius-red in histological biopsies of subcutaneous adipose tissue in a subgroup of 16 patients. Open symbols are men. **D:** Association of clinical parameters to serum  $\beta$ OHB levels in the patients studied. **E:** Patients distribution according to Fat score and biopsy-proven liver status. Numbers represent the total number of patients in stratification groups, with the number of men in parenthesis. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

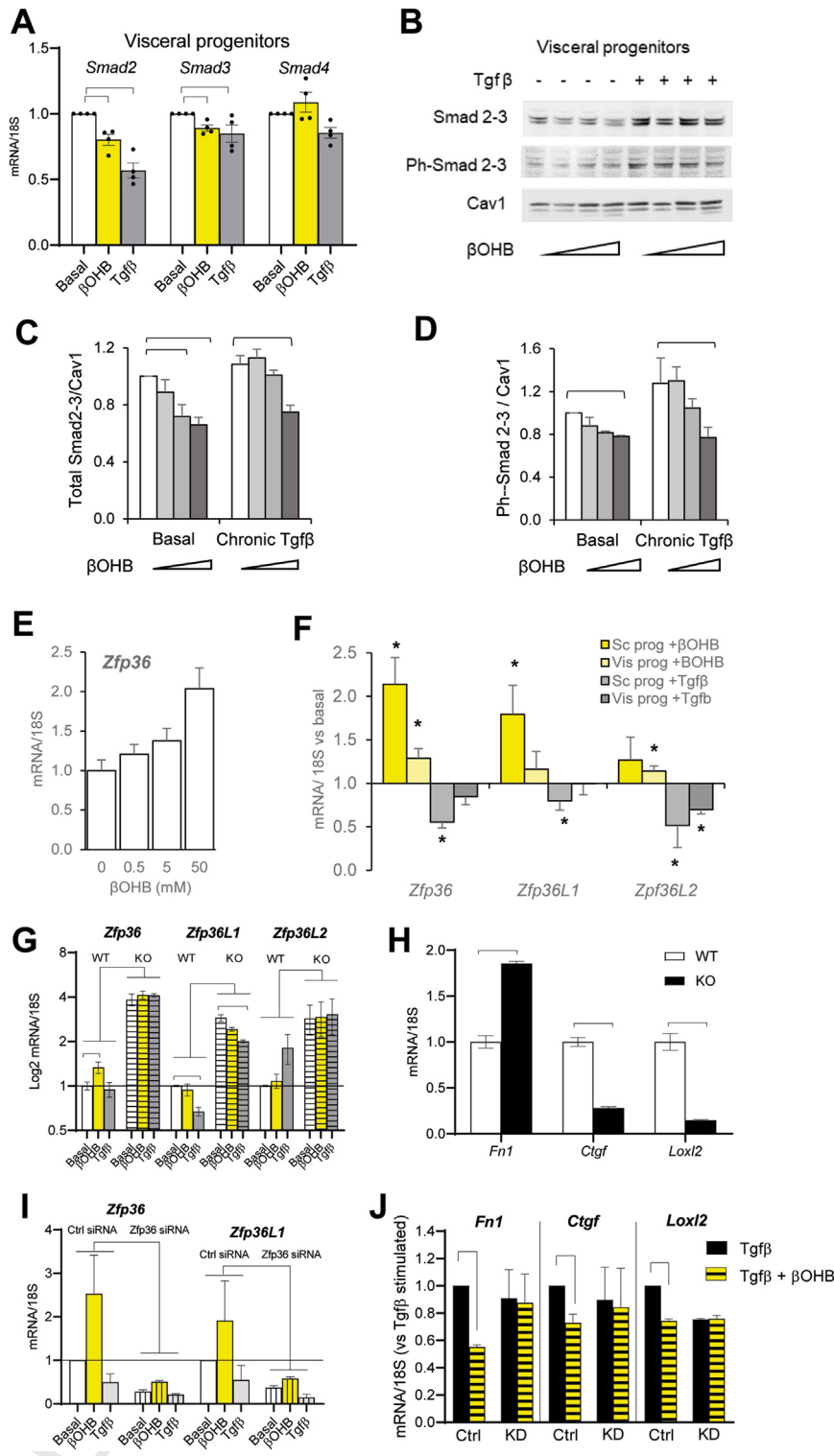
induced *Prdm16*, *Ppargc1a* and *Zfp423* when exposed to either  $\beta$ OHB or Bmp4, but displayed marginal responses in other markers. Noteworthy, in this experiment, no adipocyte differentiation cocktail was applied and progenitors were maintained for only 5 days, which might explain partial up-regulation of the browning gene program. As a whole, these data confirm the dual potential of primary cultured progenitors to respond to browning and pro-fibrotic agents, mirroring the *in vivo* situation. Noticeably, although constitutive stimulation of the PDGF receptor was shown to activate fibrotic development of progenitors in multiple mouse tissues *in vivo* [26], we could not detect pro-fibrotic responses to PDGF-A in cell culture (Suppl Fig. 2F-G), likely because PDGF receptor expression/function might be compromised by collagenase digestion or *in vitro* cell maintenance.

### 3.3. $\beta$ OHB counteracts Tgf $\beta$ stimulation of ECM genes and is anti-fibrotic

Next, progenitors were exposed to  $\beta$ OHB or Bmp4 alone for 24 h prior to addition of Tgf $\beta$  in combination with the browning agent for the following three days. We found that Tgf $\beta$  diminished expression of beige adipocyte markers in subcutaneous progenitors, alone or with  $\beta$ OHB or Bmp4 co-exposure (Figure. 2A). The anti-browning effect of Tgf $\beta$  over  $\beta$ OHB was also seen in visceral progenitors, although less marked and mostly restricted to *Tbx1* and *Ppargc1a* mRNAs (Figure. 2B). By contrast, *Ppargc1a* and *Zfp423* mRNAs were not suppressed by Tgf $\beta$  in Bmp4-stimulated visceral cells. Thus, Tgf $\beta$  selectively inhibits  $\beta$ OHB-dependent beiging.

We then examined if browning agents had the ability to counteract Tgf $\beta$ -dependent fibrotic response.  $\beta$ OHB (but not Bmp4) abolished *Ctgf* and *Loxl2* mRNA induction by Tgf $\beta$  in subcutaneous and visceral cells (Fig. 2CD-FG). *Fn1* gene induction by Tgf $\beta$  in visceral progenitors was also attenuated by  $\beta$ OHB specifically (Figure. 2E), and the  $\beta$ OHB/Tgf $\beta$  combination significantly lowered *Fn1* in subcutaneous progenitors while Tgf $\beta$  alone did not produce stimulation (Figure. 2H). *In vivo*,  $\beta$ OHB circulating concentrations change from a physiological range (0.1–0.5 mM) to more than 1 mM in fasting subjects, and rise up to 10 mM or more in diabetes.  $\beta$ OHB blunted Tgf $\beta$ -stimulated *Ctgf*, *Loxl2* and *Fn1* mRNA expression of primary progenitors in a dose dependent manner (Figure. 2I). Noticeably, in the fibroblastic 3T3-L1 cell line committed to adipogenesis, we also observed attenuated *Ctgf* mRNA induction in the combined presence of Tgf $\beta$  and  $\beta$ OHB compared to Tgf $\beta$  alone (Suppl Fig. 3A). Thus,  $\beta$ OHB can counteract pro-fibrotic action of Tgf $\beta$  in progenitors, independent of their tissue origin. Attenuation by  $\beta$ OHB of Tgf $\beta$ -dependent gene expression translated into lower Collagen1 protein contents, an effect shared with Acetoacetate, a closely  $\beta$ OHB-related ketone body (Figure. 2J).

To examine if the anti-fibrotic action of  $\beta$ OHB might have some relevance *in vivo*, we examined serum  $\beta$ OHB concentrations in a group of 30 patients with obesity involved in a bariatric surgery program, in which SCAT fibrosis was graded according to a previously defined “Fat score” integrating pericellular and perilobular tissue fibrosis, as described in [16]. Despite a similar degree of obesity in this small-sized cohort, a trend towards lower serum  $\beta$ OHB was found in those with higher Fat scores (Figure. 3A). After stratification on  $\beta$ OHB levels (median = 0.067 mM), equal distribution among highly fibrotic or non-fibrotic AT was observed in the low  $\beta$ OHB group, a proportion that fell to 25/75 in patients with high  $\beta$ OHB (Figure. 3B). Quantitative picrosirius red labelling indicated negative correlation with serum  $\beta$ OHB concentrations (Figure. 3C), consistent with an anti-fibrotic role of  $\beta$ OHB *in vivo*. Serum  $\beta$ OHB regulation obviously extends beyond AT fibrosis. In agreement, we observed that  $\beta$ OHB levels positively



**Figure 4: Mechanisms of fibrotic attenuation by  $\beta$ OHB.** **A:** Smads mRNA expression in the presence of  $\beta$ OHB or chronic Tgf $\beta$ . Bars are mean values  $\pm$  sem from 4 independent cell preparations. **B:** A representative western blot probed with antibodies against Smad2-3 (Cell signalling, ref 8685), Phospho Smad2 (Ser465/467)/Smad3 (Ser423/425) (Cell signalling, ref 8828 and Caveolin 1 (BD Transduction Laboratory, ref 610,060).  $\beta$ OHB concentration range is as in Figure 1, in the presence or absence of a chronic Tgf $\beta$  stimulation. **C-D:** Quantitative analysis of protein signal intensity after normalization with Caveolin-1 as a loading control. Bars are mean values from 3 independent cell preparations. Parentheses indicate significant differences between groups by *t* test. **E:** ZFP36 mRNA upregulation by  $\beta$ OHB. **F:** Expression of ZFP36 and related transcripts by  $\beta$ OHB and Tgf $\beta$  in subcutaneous and visceral progenitors. Bars are mean values  $\pm$  sem from 3 to 5 independent cell preparations. \* indicate significant differences compared to basal by Student *t* test. **G-H:** Gene expression in fibroblast cell lines from Wild Type (WT) or ZFP36 KO mice. Note that in KO mice, ZFP36 mRNA is transcribed but contains an insertion in exon 2 which prevents protein production. **I-J:** Knock-down of ZFP36 with siRNA in primary progenitors maintained with Tgf $\beta$  with or without  $\beta$ OHB. Bars are mean values from 3 independent experiments. Parentheses indicate significant differences between groups by *t* test.



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1 correlated with serum free fatty acids or systolic blood pressure, and  
2 negatively associated to triglyceridemia and thyroid stimulating hor-  
3 mone (Figure. 3D). The results were not influenced by gender.  
4 Regarding liver status that could be documented from a liver biopsy in  
5 25 patients of the cohort, no link was found between Fat score and  
6 liver fibrosis. An equal proportion of adipose tissue samples were  
7 scored for high or low fat fibrosis among patients with normal liver  
8 histology or non-alcoholic fatty liver disease (NAFLD). Even, in 10  
9 patients with biopsy-proven non-alcoholic steato-hepatitis (NASH),  
10 only 2 had elevated Fat score, whereas 8 had non-fibrotic fat tissue  
11 (Figure. 3E).

### 3.4. Mechanisms of $\beta$ OHButyrate anti-fibrotic activity by canonical smad signalling and non-canonical Zfp36 induction

#### 3.4.1. $\beta$ OHButyrate reduces smad activation in visceral progenitors

12  $\beta$ OHB is metabolized into acetylCoA that enter the TCA cycle. We  
13 reasoned that if  $\beta$ OHB metabolism was involved, its impact should be  
14 stronger in conditions of carbon sources limitation. Progenitors were  
15 cultured in Gln High, a high glucose, glutamine-supplemented me-  
16 dium, or in Gln Low, a high glucose, glutamine-depleted medium, in  
17 which glutamine was only available from serum. Such reduced  
18 glutamine/high glucose conditions the AT microenvironment found in  
19 obese diabetic subjects, as described [27]. We observed that  $\beta$ OHB  
20 equally attenuated Tgf $\beta$ -dependent elevation of *Ctgf* mRNA in low or  
21 high Gln (Suppl Fig. 3B), suggesting that modulation of carbon entry in  
22 the TCA cycle is not a major contributor of  $\beta$ OHB anti-fibrotic effect.  
23 Tgf $\beta$  regulates the expression of ECM genes by activation of SMAD-2/3  
24 downstream of Tgf $\beta$  receptors [28]. We first considered that anti-  
25 fibrotic  $\beta$ OHB might act at the pre-receptor level by production of  
26 extracellular modulators able to restrain ligand binding or presentation  
27 [29]. However, exposure to a *bona fide* exogenous Tgf $\beta$  antagonist,  
28 follistatin, alone or in combination with  $\beta$ OHB, had no effect on visceral  
29 progenitors ECM gene expression (Suppl Fig. 3C). We next examined  
30 SMAD-2/3 regulation, whose phosphorylation on serine residues  
31 423–426 is induced downstream of Tgf $\beta$  receptor. We observed that  
32 *Smad2* and *Smad3* mRNAs were slightly but significantly reduced by  
33  $\beta$ OHB treatment in, whereas *Smad4* mRNA did not change (Figure.  
34 4A). Western blots confirmed a dose-dependent decrease (30–40%)  
35 of Smad-2/3 protein contents upon  $\beta$ OHB exposure, in both unstim-  
36 ulated cells or after chronic Tgf $\beta$  stimulation (Figure 4B,C). Moreover,  
37 SMAD-2/3 serine phosphorylation was reduced in chronically Tgf $\beta$   
38 treated visceral progenitors (Figure. 4D). Thus, blunting of Smad-2/3  
39 dependent Tgf $\beta$  signalling by  $\beta$ OHB likely contributes its anti-fibrotic  
40 response. However, in similar experiments with subcutaneous pro-  
41 genitors, Smad-dependent Tgf $\beta$  response was not significantly  
42 affected by  $\beta$ OHB (data not shown), suggesting additional mechanisms  
43 might operate depending on cell origin.

#### 3.4.2. Identification of ZFP36 as a mediator of $\beta$ OHB anti-fibrotic action

44 To identify other  $\beta$ OHB-driven mechanisms, we explored data  
45 generated from single cell RNA sequencing studies, which document  
46 specific AT stromal gene expression in progenitor subgroups. From  
47 publicly available reports [19–21], data sets detailing information from  
48 chow fed mice at different developmental states, AT anatomical lo-  
49 cations or sex were downloaded, integrated using Batch Balanced K-  
50 nearest Neighborhood algorithm (<https://doi.org/10.1093/bioinformatics/btz625>), and processed by Leiden clustering for community  
51 detection (<https://doi.org/10.1038/s41598-019-41695-z>), which  
52 defined groups with similarity relationships. We found that *Fabp4*,

53 *Cd36*, *Pparg* and *Cav1* genes, associated with adipogenesis, top  
54 ranked in three clusters, likely preadipocyte subsets at different stages  
55 (Suppl Fig. 4A). Using the ranked gene list to define group identity, we  
56 preselected a cluster comprising top ranking expression of ECM related  
57 genes (*Col1a1*, *Col3a1*, *Gsn*, *Dcn*) and the progenitor gene marker  
58 *Pdgfra*. This progenitor cluster with high potential for ECM deposition  
59 also contained Tgf $\beta$  receptor genes, suggesting responsiveness to  
60 pro-fibrotic factors. A top ranked gene in this cluster was *ZFP36*, which  
61 encodes a protein known as Tristetraprolin (TTP), a member of the  
62 “early response gene” family closely related to *ZFP36L1* and *ZFP36L2*.  
63 *ZFP36* proteins belong to a family of RNA binding proteins known to  
64 destabilize AU-rich element (ARE)-containing mRNAs by promoting  
65 the removal of their polyA tails [30]. Indeed, *ZFP36* was found to  
66 decrease the stability of anti-adipogenic *Tnfa* mRNA, and *ZFP36L1*  
67 could target inflammatory transcripts of numerous components of  
68 senescence-associated secretory phenotype.

69 Search in ARED-Plus (<http://brp.kfshrc.edu.sa/ared>), the AU-Rich  
70 Element Database [31] revealed hits in the 3'UTRs of *Ctfg* and *Fn1*  
71 mRNAs, making *ZFP36* an attractive candidate regulator of ECM  
72 transcript stability. Moreover, a survey of *ZFP36* expression patterns  
73 from public databases indicated preferential expression of *ZFP36*  
74 mRNA, and related *ZFP36I1* and *ZFP36I2* in BAT versus white fat (GSE  
75 8044), while in GSE40486, BAT *ZFP36* was induced by cold (Suppl  
76 Fig. 4B). High-throughput studies in T cells highly expressing *ZFP36*,  
77 to identify direct targets by sequencing UV crosslinked immunopre-  
78 cipitated transcripts, revealed hundreds of mRNAs to which *ZFP36*  
79 could bind [32], among which Glutaminase (encoded by *Gls* gene).  
80 Functionally, glutaminolysis is a pathway that feeds proline synthesis,  
81 a highly abundant amino acid in collagens. Thus, links from gluta-  
82 minase to collagen deposition through *ZFP36* has functional relevance.  
83 We found that Tgf $\beta$  stimulated *Gls* expression more than 3-fold in 3T3-  
84 L1 preadipocytes, an effect that was completely blunted by the  $\beta$ OHB/  
85 Tgf $\beta$  combination (Suppl Fig. 3D). Such a similarity in the regulation of  
86 *Gls*, a *bona fide* *ZFP36* target, with that of *Ctgf*, and *Fn1*, bearing  
87 potential *ZFP36* binding sequences, reinforced the possibility that  
88 these might be under *ZFP36* post-transcriptional control.

89 Consistent with a role for *ZFP36* in  $\beta$ OHB anti-fibrogenic effects, we  
90 found that primary subcutaneous progenitors *ZFP36* mRNA dose-  
91 dependently increased (2-fold) upon  $\beta$ OHB exposure (Figure. 4E).  
92  $\beta$ OHB also stimulated the expression of the closely related *ZFP36I1*  
93 and *ZFP36I2* (Figure. 4F). Conversely, pro-fibrotic stimulation with  
94 Tgf $\beta$ , in which optimal stability of ECM transcripts is expected, led to a  
95 significant drop in *ZFP36* and related *ZFP36I1* and 2 (Figure. 4F).  
96 Comparable regulation by  $\beta$ OHB and Tgf $\beta$  was observed in visceral  
97 progenitors, although with lesser magnitude (Figure. 4F).

98 We next investigated mouse fibroblast cell lines derived from *ZFP36*  
99 null (KO) or control (WT) mice [17]. Knock-out of *ZFP36* expression [33]  
100 was achieved by insertion of a neomycin cassette into the second exon  
101 of the *ZFP36* gene, leading to the synthesis of a non-functional *ZFP36*  
102 *fusion* mRNA detected at high levels in KO compared to WT cells  
103 (Figure. 4G), in accordance with previously reported *ZFP36* autor-  
104 egulation. In favour of a direct control of *Fn1* mRNA abundance by  
105 *ZFP36*, *Fn1* was upregulated in KO versus WT cells (Figure. 4H).  
106 However, *Ctgf* and *Loxl2* mRNAs were strongly suppressed in the KO  
107 fibroblasts (Figure. 4H).

108 We then performed *ZFP36* knock-down in primary subcutaneous  
109 progenitors using siRNAs which also down-regulated *ZFP36I1* (Figure.  
110 4I).  $\beta$ OHB was no longer able to restrain Tgf $\beta$ -mediated stimulation of  
111 *Ctgf*, *Loxl2* and *Fn1* in *ZFP36/ZFP36L1* deficient progenitors (Figure.  
112 4J). Thus, *ZFP36* expression is required for  $\beta$ OHB anti-fibrotic ac-  
113 tion. However, the ability of  $\beta$ OHB to stimulate genes associated with  
114

1 beige adipogenesis was not affected in *ZFP36* deficient progenitors  
2 (Data not shown).

#### 4. DISCUSSION

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6 In response to nutritional and environmental cues, fat tissue is  
7 constantly remodeled, a process involving adipose progenitors dialog  
8 with mature adipocytes and other immune cell actors, to modulate  
9 adipogenesis or tissue fibro-inflammation. Our study documents a  
10 reciprocal cross talk between brown-like adipogenesis and fibrogenic  
11 activation of progenitors. We show that metabolites promoting brown-  
12 like fat cell development might also alleviate AT responses to profi-  
13 brotic Tgf $\beta$ , suggesting potential targeted approaches to reduce AT  
14 fibrosis development. First, mirroring previously reported improvement  
15 of metabolic health by adipose tissue thermogenic activation in rodents  
16 [5,6], we show here that raising mice at thermoneutrality to induce  
17 thermogenic dormancy further aggravates AT fibrotic response to HFD.  
18 We demonstrate that  $\beta$ -hydroxybutyrate, a metabolite able to activate  
19 brown-like adipogenesis, can also counteract Tgf $\beta$ -dependent pro-  
20 fibrotic responses in primary cultured progenitors. We identified  
21 different pathways that underline this cross talk, governed by tissue  
22 origin of adipose progenitors (visceral or subcutaneous). In visceral  
23 progenitors,  $\beta$ OHB downregulates the expression and phosphorylation  
24 of Smads downstream of Tgf $\beta$  receptor activation, whereas in sub-  
25 cutaneous preadipocytes,  $\beta$ OHB is found to promote post-  
26 transcriptional regulation of Tgf $\beta$  gene targets notably through in-  
27 duction of *ZFP36*/Tristetraprolin, a protein regulating mRNA destabi-  
28 lisation. The mechanism by which  $\beta$ OHB exerts its anti-fibrotic effect  
29 might be accounted for its property as a signalling molecule or a  
30 metabolic substrate. As such, intracellular import of ketone bodies by  
31 monocarboxylate transporters is shared with that of lactate, and lactate  
32 fluxes are important players in beige adipocyte phenotype [34].

33 Importantly, our *in vitro* observations may have translational relevance  
34 since circulating  $\beta$ -hydroxybutyrate levels in patients with obesity  
35 inversely associate with subcutaneous adipose tissue fibrosis.  $\beta$ -  
36 hydroxybutyrate is a circulating metabolite that increases during  
37 ketogenesis [35], a fasting-dependent pathway. It is mainly produced  
38 by the liver as a product of partial fatty acids oxidation; however,  
39 adipose tissue was recently established as an alternative source. In  
40 particular, mature adipocytes were found to produce  $\beta$ -hydroxy-  
41 butyrate within the AT local microenvironment, which was shown to  
42 directly impact tissue progenitors towards brown-like fat cell differ-  
43 entiation [23]. Importantly, we show here that in the presence of Tgf $\beta$ ,  
44 although  $\beta$ OHB was ineffective in maintaining brown-like fat cell  
45 orientation (Figure. 2AB), it could efficiently counteract the induction  
46 of ECM genes. This suggests that raising  $\beta$ OHB in the obese AT context  
47 might have some interest against fibrosis development. Interestingly, a  
48 new class of anti-diabetic drugs designed to promote glucose renal  
49 excretion, referred to as “SGLT2 inhibitors” were shown to increase  $\beta$ -  
50 hydroxybutyrate levels in mouse models [36]. The possibility that  
51 beneficial metabolic effects of these drugs could be partly mediated by  
52 beta-hydroxybutyrate-dependent remodeling of adipose tissue is an  
53 interesting perspective that remains to be explored.

54 We identified the RNA binding protein *ZFP36*/Tristetraprolin as a  
55 mediator of the anti-fibrotic effect of  $\beta$ -hydroxybutyrate. *ZFP36* is  
56 considered a broad anti-inflammatory molecule, acting on post-  
57 transcriptional regulation of many immune-related genes by decreasing  
58 the stability of corresponding mRNAs [37–39]. *ZFP36*/  
59 Tristetraprolin null mice suffer from hyper-inflammation associated  
60 with cachexia, arthritis and dermatitis, which can be prevented by  
61 treatment with anti-TNF $\alpha$  antibodies [33]. Conversely, it was also

demonstrated that increased Tristetraprolin expression could protect  
mice against immune-mediated inflammatory pathologies [40]. We  
report the induction of *ZFP36* gene expression by  $\beta$ OHB, and its  
downregulation by Tgf $\beta$ , along with changes in Tgf $\beta$ - or  $\beta$ OHB-  
mediated fibrotic responses of progenitors. The present study extends  
the spectrum of *ZFP36* targets to several Tgf $\beta$ -regulated mRNAs.  
Among those, *Ctgf* induced by the canonical Tgf- $\beta$ /Smad pathway [41],  
is a crucial downstream effector in mediating Tgf $\beta$ -related fibro-  
genesis [42]. Its inhibition might not only prevent but also reverse  
fibrosis progression [43]. Based on its broad actions, *ZFP36* has great  
potential to impact the progenitor niche towards attenuation of ECM  
gene program and concomitant reduction in inflammation [44]. In  
accordance with the present findings that *ZFP36* expression dampens  
adipose progenitor response to profibrotic Tgf $\beta$ , several reports have  
associated *ZFP36*/Tristetraprolin deficiency with liver fibrosis aggra-  
vation [45], or its over-expression to protection against lung fibrosis  
following ischemia/reperfusion [46]. In the colonic epithelium, *ZFP36*  
was also shown to target *iNos* mRNA [47], an important mediator in  
fibro-inflammation regulated by Tnf $\alpha$ . Recently, the turnover of the  
peptide hormone FGF21 was found to be directly regulated by *ZFP36*  
family members in the context of alcoholic [48] or non-alcoholic hep-  
atic steatosis [49], which extends the spectrum of *ZFP36* targets to  
metabolic regulation. Thus, the *ZFP36* family has great potential as a  
multi-faced post-transcriptional regulator coordinating inflammation,  
extracellular matrix composition and metabolic responses.

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#### APPENDIX A. SUPPLEMENTARY DATA

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.molmet.2022.101512>.

#### CONFLICT OF INTEREST

None declared.

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