Acute induction of uncoupling protein 1 by citrulline in cultured explants of white adipose tissue from lean and high-fat-diet-fed rats

Nolwenn Joffin¹,², Anne-Marie Jaubert²,³, Jessica Bamba¹,², Robert Barouki¹,², Philippe Noirez¹,⁴,⁵, and Claude Forest¹,²,*

¹Université Paris Descartes, Sorbonne Paris Cité; Paris, France; ²Institut National de la Santé et de la Recherche Médicale UMR-S 1124; Faculté des Sciences Fondamentales et Biomédicales; Pharmacologie Toxicologie et Signalisation Cellulaire; Paris, France; ³Département de Biochimie et de Biologie Moléculaire; Faculté de Médecine Paris-Ile de France-Ouest; Université de Versailles Saint-Quentin en Yvelines; Versailles, France; ⁴Institut de Recherche Biomédicale et d’Épidémiologie du Sport, EA 7329; Paris, France;
⁵Faculté des Sciences et Techniques des Activités Physiques et Sportives; Paris, France

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A diet enriched with citrulline (CIT) reduces white adipose tissue (WAT) mass. We recently showed that CIT stimulated β-oxidation in rat WAT explants from young (2–4 months) but not old (25 months) rats. Here we show that both in old rats and high-fat-diet-fed young rats, uncoupling protein one (UCP1) mRNA and protein expressions were weaker than those in young control rats. Selectively in WAT from young rats, a 24h CIT treatment up-regulated expressions of UCP1, peroxisome proliferator-activated receptor-α (PPARα), PPARγ co-activator-1-α and mitochondrial-transcription-factor-A whereas it down-regulated PPARγ2 gene expression, whatever the diet. These results suggest that CIT induces a new metabolic status in WAT, with increased β-oxidation and uncoupling of respiratory chain, resulting in energy expenditure that favors fat mass reduction.

Introduction

Excess energy storage in white adipose tissue (WAT) favors an increase in fat mass with pathophysiological consequences like insulin resistance and altered glucose tolerance leading to type-2 diabetes (T2DM). Aging and high-fat diet (HFD) often result in enlarged WAT mass, particularly intra-abdominal depots like the retroperitoneal (RET). One of the strategies to oppose fat mass gain would be to activate energy dissipation in brown adipose tissue (BAT). Brown adipocytes have the unique feature to express the uncoupling protein 1 (UCP1), responsible for the uncoupling between oxidative phosphorylation and ATP production, therefore dissipating energy as heat.¹,² In 1992, Cousin et al. demonstrated the presence of UCP1-expressing adipocytes in well-defined WAT depots.³ Such cells, resembling brown adipocytes in WAT, were further described and named BRITE for “brown-in-white” or beige adipocytes, then shown as having an intermediate signature between brown and white adipocytes.⁴,⁵ Although the distinct origins of these cells remain a matter of debate, they both express UCP1.⁶,⁷ Activators of UCP1 in BAT and inducers of WAT browning, i.e. the acquisition of a brown phenotype, have been determined. Among these agents, cold exposure, β-adrenergic agonists, retinoids, triiodothyronine, irisin, atrial natriuretic factor, prostaglandins, adenosine or lipolytic NEFA themselves have been shown to up-regulate UCP1.⁸,¹¹

An interesting observation of decreased WAT mass together with increased BAT mass and UCP1 arose from high-fat-diet-induced (HFD) overweight rats to which L-arginine (L-ARG) was given in drinking water.¹² L-ARG induced lipolysis and β-oxidation in WAT.¹³ These studies were carried out either in vivo or in cultured cells. Studies on cultured explants from WAT are scarce. Using RET explants, we showed recently that citrulline (CIT) induced lipolysis.
and β-oxidation while it reduced glyceroneogenesis and NEFA re-esterification in control and HFD-fed young rats. Similar data were obtained on RET explants from old rats, except that β-oxidation remained unaffected by CIT. CIT is an amino acid that is not used for protein synthesis. Like L-ARG, CIT is an intermediary product in nitric oxide (NO) synthesis, provided that argininosuccinate synthase (ASS) and argininosuccinate lyase (ASL) are present, which is the case in macrophages but not in isolated adipocytes. Unlike what occurs with L-ARG, CIT, administered as a food supplement, is not metabolized in the liver, therefore its biodisponibility is higher than that of L-ARG. CIT is thus proposed as a food supplement during aging and in sports to improve muscle mass and function.

In the present study we investigated whether CIT exerted a browning effect on epididymal (EPI) and RET WAT explants from control and HFD-fed young rats together with old rats. To this aim we analyzed variations in UCP1 expression and in its main transcriptional regulators.

**Results**

**Effect of age, diet, and CIT on the expression of UCP1 gene and protein**

We first evaluated the effect of age or HFD on the relative UCP1 gene and protein expressions in RET and EPI WAT explants maintained in culture for either 8h or 24h. For CD in both depots, UCP1 mRNA level was similar in explants from 2-month-old and 4-month-old rats while for HFD, it presented a 45–55% decrease (Fig. 1A). In both depots from 25-month-old rats, UCP1 mRNA and protein levels were significantly lower than in explants from younger rats. Treatment with CIT did not significantly influence the expression of UCP1 in control rats (Fig. 1C). However, in HFD-fed rats, CIT treatment increased the expression of UCP1 mRNA and protein (Fig. 1D).

![Figure 1](image-url)
old rats, UCP1 mRNA was reduced to 50–60% when compared to 2-month-old rats (Fig. 1A). A similar pattern of reduction of UCP1 protein occurred with 40–50% decrease at 24h of treatment (Fig. 1B). We then assessed the effect of CIT treatment for 8h and 24h on UCP1 expression in RET and EPI WAT explants. At 24h CIT treatment, UCP1 mRNA was increased 3.5 to 6-fold, for both depots, in 2-month-old and 4-month-old rats, whether the latter were maintained or not on a HFD diet for 8 weeks, while it remained unaffected in WAT from 25-month-old rats (Fig. 1C). A weaker (2.3-fold) increase in UCP1 mRNA was observed at 8h CIT treatment in EPI WAT from 2-month-old rats, although no other change occurred at this time (Fig. 1C).

At 24h CIT exposure, UCP1 protein was upregulated 1.5- to 2-fold in WAT from young rats for both depots, whatever the diet, while it remained unaffected in WAT from 25-month-old rats (Fig. 1D).

**Effect of CIT on the expression of genes encoding transcription factors and regulators involved in mitochondria metabolism**

In agreement with our previously published results, peroxisome proliferator-activated receptor α (PPARα) gene expression was stimulated by CIT in RET explants from 2-month-old young rats whereas it remained unaffected in RET explants from 25-month-old rats (Fig. 2A and B).14 In contrast, PPARγ2 gene expression was downregulated (58–86%) by CIT in RET explants whatever the age of the animals (Fig. 2A and B).14 This reduced level was observed at 8h of exposure to CIT and remained stable at 24h. These previously described observations were extended to RET explants from CD-fed and HFD-fed 4-month-old rats that responded in a manner similar to those from 2-month-old rats (Fig. 2A and B). In addition, the response of EPI explants to either...
8h or 24h of CIT exposure was similar to that of RET, except for 25-month-old rats in which the PPARγ2 gene remained unaffected (Fig. 2C and D). CIT treatment augmented PPARγ co-activator 1α (PGC-1α) and mitochondrial transcription factor A (TFAM) mRNA in both depots from young rats whatever the diet (Fig. 2A–D). These increases were more pronounced at 24h than at 8h of treatment. In explants from 25-month-old rats, PGC-1α and TFAM genes remained unresponsive to CIT whatever the WAT depot and the time of treatment (Fig. 2A–D).

**Discussion**

We show that UCP1 mRNA and protein are expressed in RET and EPI WAT from rats, in accordance with previously published studies. This expression is reduced with either aging or HFD. Our previous results established that CIT induced direct, rapid and deep modifications of NEFA metabolism in WAT explants from rats. Notably, CIT increased β-oxidation capacity, together with the expression of the PPARα gene, in RET explants from young rats whether they were fed a CD or a HFD. Here, we observe that CIT was also able to induce UCP1 in WAT from young animals. However, no change in β-oxidation, UCP1 expression and its transcriptional regulators (the present study) occurred in explants from old animals in response to this amino acid. The mechanism that underlines this young versus old difference in β-oxidation and UCP1 expression remains an open issue.

Therefore, in old animals, part of lipolytic NEFA are released from WAT and can be used as energetic substrates for demanding tissues like muscle. Such observations are in line with the increase in muscle mass from old rats fed with either a L-ARG- or a CIT-supplemented diet. We observed previously that in young animals, whether HFD-fed or not, NEFA output remained unaffected by CIT but NEFA oxidation was stimulated. In the present study, we show that CIT up-regulates UCP1 expression in WAT explant from young rats. This effect is rapid since a 3-fold to 6-fold increase in UCP1 mRNA occurred at 24h of treatment, together with a 1.5- to 2-fold rise in the protein. Such a new metabolic feature is the sign of a situation of energetic expenditure in WAT.

What could be the mechanism of CIT action? Since cAMP, through protein kinase A (PKA) activation, is one of the major inducers of UCP1 gene expression in brown adipocytes, it could be the mediator of CIT action, although there is no report of the effect of this amino acid on PKA. Such a mechanism is likely not occurring here. One of the best-known cAMP-responsive gene in adipocytes is pck1 (the cytosolic phosphoenolpyruvate carboxykinase—PEPCK-C—gene). Therefore, any stimulation by CIT of cAMP production should induce pck1 but the inverse occurs here. However, CIT stimulates HSL phosphorylation in RET WAT and induces NEFA release when explants are isolated from HFD-fed young rats or from old rats. Among the other agents that were demonstrated as capable of inducing HSL phosphorylation and lipolysis, the mitogen-activated protein kinase pathway and extracellular signal-regulated kinase (ERK) could play a role. Alternatively, lipolysis could be activated because of the reduction of an inhibitory pathway, like those mediated by the AMP-activated protein kinase (AMPK), calcium/calmodulin-dependent protein kinase II or glycogen synthase kinase IV. Interestingly, L-ARG was reported as being both lipolytic and capable of inducing AMPK, potentially opposite phenomena in adipocytes, without clear explanations for this apparent contradiction.

One of the best candidates for mediating CIT action could be NO. As mentioned above, ASS and ASL are not expressed in WAT adipocytes. Hence, any NO production from CIT would primarily occur in macrophages present in WAT and would diffuse from macrophages to adipocytes where it would act on UCP1 expression. Previous reports showed the lipolytic and anti-glyceroenogenic nature of NO in WAT. Besides, NO induces mitochondrial biogenesis via guanosine 3',5'-monophosphate (cGMP) and PGC-1α, the central transcriptional activator in thermogenic adipocytes. Both PPARα and PPARγ are transcription factors that interact with PGC-1α. CIT strongly induced PGC1α and PPARα gene expressions in both RET and EPI WAT from young rats, whereas PPARγ2 mRNA was down-regulated in the same animals, in agreement with our previous results. The reduction in PPARγ2 reflects the previously observed strong decrease in PEPCK-C expression and in glyceroenogenesis. The simultaneous induction by CIT of PGC1α and PPARα is in line with the observed concomitant up-regulation of UCP1 in WAT, probably leading to a rise in the thermogenic status of the cells. CIT effect on UCP1 gene expression is supposedly transcriptional through a PPARα/PGC1α-process, since this couple was clearly demonstrated as acting as an inducer of the UCP1 gene in rodents and in humans. PPARα and PGC1α were also shown to stimulate FA oxidation, notably in brown adipocytes, via the induction of expression of CPT 1b and VLCAD. These observations are in agreement with our previous results showing that selectively in WAT explants from young rats, CIT induced an increase in β-oxidation capacity, together with an upregulation of CPT 1b and VLCAD.

Another interesting observation of the present study is the CIT effect on TFAM expression because TFAM is a PGC1-α target and a major transcription factor for mitochondrial DNA replication. The possible CIT action as a factor favoring mitochondriogenesis during WAT browning remains an open issue with potential physiological consequences. Such a question also occurs with L-ARG-fed rats for which PGC 1α and mitochondriogenesis are enhanced, together with BAT and whole-body energy metabolism.

Altogether, our data identify a novel mechanism of CIT health effects and highlight the induction of UCP1 as a mediator of decreased WAT mass, browning of WAT and increased energy expenditure. The data open new and simple avenues of therapeutic intervention in overweight and obesity.
Materials and Methods

Animals: housing and feeding

Two-month-old male Sprague Dawley rats from Center d’Elevage de Rats Janvier (Berthevin, France), were fed ad libitum with a standard (control) diet (CD) (n = 21). They were acclimatized for 2 weeks then randomized into 3 groups. The two-month-old CD rats (n = 7) were immediately used for the culture of AT explants. The two other groups were maintained for an additional 8 weeks under either a CD (4-month-old CD; n = 7) or a high-fat diet (4-month-old HFD; n = 7). Twenty-five-month-old male Sprague Dawley rats, fed ad libitum with CD, were a gift from Sainte Anne Hospital (Paris, France) (n = 7). All rats have free access to water and food ad libitum and were housed at constant room temperature (24°C) on a 12-h-light/dark cycle. CD rats were fed with a standard balanced diet (11.7 MJ/kg) (64% carbohydrates, 20% proteins, 8% lipids, 8% vitamins and minerals) from Safe (Augy, France). HFD rats were fed with a HFD (19.5 MJ/kg) (64% carbohydrates, 20% proteins, 3% lipids, 8% vitamins and minerals) from Safe (Augy, France). Rats were anaesthetized and after blood sampling from the jugular vein, they were killed by decapitation. Samples of WAT were removed and weighted. The protocol for the animal studies was conducted according to the French Guidelines for the care and use of experimental animals.

Rat adipose tissue explants

EPI and RET WAT from rats were sampled for analyses. About 200mg of tissue were cut in about 20mg fragments and cultured in Krebs Ringer buffer saline (KREBS) medium as previously described.14,15 Explants were exposed to 2.5 mmol/L CIT for 8h or 24h then frozen before further studies.

RNA extraction, cDNA synthesis and real-time RT-PCR

Total RNA was extracted from RET WAT with Rneasy lipid tissue mini kit from Qiagen. RNA was quantified by spectrophotometric Nanodrop. Total RNA (500ng) was reverse transcribed using SuperScript® III First-Strand Synthesis SuperMix kit from Life Technologies. Samples of cDNA were diluted 1:40, amplified and used for RT-qPCR measurements using SYBR Green from Roche as previously described.14 RT–qPCR was performed in the LightCycler1536 instrument from Roche as follows: 75°C for 2min, 95°C for 10min, 40 cycles of denaturation (95°C for 15s to the denaturation, 60°C for 30s to the annealing, 72°C for 30s to the extension). Results were analyzed with the LightCycler1536 software from Roche. Ribosomal RPL13 RNA was used to normalize cDNA. Quantification of mRNA was carried out by comparing the number of cycles required to reach reference and target threshold values (2^-ΔΔCt) method as described previously.43 Sequences of the rat sense and antisense nucleotides were: PGC1-α : 5’-tgctacggttgaagaggc-3’ 5’-gcaggtggagctgtctg-3’; PPARα : 5’-aacggctctctgctgtagc-3’ 5’-cgagtttgcagcag-3’; PPARγ : 5’-ttgctcggttagggtgaa-3’ 5’-caagagatggagtctg-3’; TFAM : 5’-gctccggcggagtggtgg-3’ 5’-ggacggaggttccg-3’; UCP1 : 5’-tataggttatagcaccaca-3’ 5’-tgaaagctcattctgtgg-3’; RPL13 : 5’-tgccagggctcag-3’ 5’-tgccatcacggtc-3’. Western blot

Protein fractions were prepared from WAT explants in a buffer containing 10mmol/L HEPES ph 7.9, then separated by SDS-PAGE and transferred to nitrocelullose filters. Blots were blocked with 3% BSA in PBS containing 0.05% Tween 20 and incubated with 1:1000 and 1:500 dilutions for the UCP1 and β-actin antisera, respectively. Rabbit UCP1 and mouse β-actin antisera were purchased from thermofisher and Abcam respectively.

The Odyssey method was used for detection, following the procedure described by the manufacturer (Li-COR). Quantitative results of Western blotting were obtained by densitometry in ImageJ software.

Statistical analysis

Data are presented as means ± SEM. Each independent experiment was carried out in triplicate. Statistical analysis was carried out using the nonparametric Mann & Whitney U test for pairwise comparisons, which was applied due to the small number of experiments (n < 10). A value of P < 0.05 was considered as significant.

* P < 0.05 vs. control; ** P < 0.01 vs. control

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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