Nutritional, hormonal, and depot-dependent regulation of the expression of the small GTPase Rab18 in rodent adipose tissue

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Abstract

There is increasing evidence that proteins associated with lipid droplets (LDs) play a key role in the coordination of lipid storage and mobilization in adipocytes. The small GTPase, RAB18, has been recently identified as a novel component of the protein coat of LDs and proposed to play a role in both β-adrenergic stimulation of lipolysis and insulin-induced lipogenesis in 3T3-L1 adipocytes. In order to better understand the role of Rab18 in the regulation of lipid metabolism in adipocytes, we evaluated the effects of age, fat location, metabolic status, and hormonal milieu on Rab18 expression in rodent white adipose tissue (WAT). Rab18 mRNA was undetectable at postnatal day 15 (P15), but reached adult levels by P45, in both male and female rats. In adult rats, Rab18 immunolocalized around LDs, as well as within the cytoplasm of mature adipocytes. A weak Rab18 signal was also detected in the stromal-vascular fraction of WAT. Rab18 mRNA was undetectable at postnatal day 15 (P15), but reached adult levels by P45, in both male and female rats. In adult rats, Rab18 immunolocalized around LDs, as well as within the cytoplasm of mature adipocytes. A weak Rab18 signal was also detected in the stromal-vascular fraction of WAT. In mice, fasting significantly increased, though with a distinct time-course pattern, Rab18 mRNA and protein levels in visceral and subcutaneous WAT. The expression of Rab18 was also increased in visceral and subcutaneous WAT of obese mice (diet-induced, ob/ob, and New Zealand obese mice) compared with lean controls. Rab18 expression in rats was unaltered by castration, adrenalectomy, or GH deficiency but was increased by hypophysectomy, as well as hypothyroidism. When viewed together, our results suggest the participation of Rab18 in the regulation of lipid processing in adipose tissue under both normal and pathological conditions.

Key Words
► fasting
► gene expression
► lipid metabolism
► obesity
► Rab18
► white adipose tissue

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Introduction

Adipose tissue has received much attention over the last two decades due to the discovery of its role as an active endocrine organ (Kershaw & Flier 2004). Moreover, recent discoveries have also significantly changed the classical view of adipose tissue as an inert energy depot and unveiled the complex regulation of adipose tissue lipolysis and lipogenesis. Indeed, regulation of lipid metabolism in adipocytes depends on a fine balance between multiple factors, both extracellular and intracellular (Brasaemle 2007, Koldtiz & Langin 2010, Watt & Spriet 2010). Although to date there exists a wealth of information on the extracellular factors that regulate adipocyte function, the intracellular components and mechanisms that participate, directly or indirectly, in the control of lipid trafficking and metabolism are less well known.

Lipid droplets (LDs) are responsible for the storage of excess energy in the form of neutral lipids and thus play a key role in adipocytes (Murphy 2001, Wolins et al. 2006). LDs are coated by proteins that are required for optimal lipid storage and fatty acid release (Brown 2001, Goodman 2008). LD coat proteins include lipid metabolic enzymes (Brasaemle et al. 2004, Kuerschner et al. 2008, Miyoshi et al. 2008, Guo et al. 2009), the PAT family of proteins (named after perilipin, adipophilin, and TIP47) that coordinate lipid storage and utilization (Brasaemle 2007, Ducharme & Bickel 2008), the cell death-inducing DFF45-like effector (CIDE) family of proteins that regulate lipogenesis (Gong et al. 2009), and proteins regulating membrane trafficking events, which include RAB and ARF GTPases, soluble N-ethylmaleimide-sensitive factor activating protein receptor (SNARE) proteins, and motor proteins (Liu et al. 2007, Guo et al. 2008, Hommel et al. 2010). In particular, several members of the RAB family of small GTPases have been shown to decorate the surface of LDs (Brasaemle et al. 2004, Murphy et al. 2009), yet the precise function of most RAB proteins expressed in adipocytes remains unknown. Among those best documented is RAB18, which was first shown to associate with the surface of LDs in 3T3-L1 adipocytes in response to β-adrenergic stimulation and was therefore proposed to be involved in the release of lipids from LDs (Brasaemle et al. 2004, Martin et al. 2005, Ozeki et al. 2005). Consistent with these early studies, we recently demonstrated that RAB18 overexpression increased basal lipolysis and RAB18 silencing-impaired forskolin-stimulated lipolysis in 3T3-L1 cells (Pulido et al. 2011). Intriguingly, we also observed that insulin, like the β-adrenergic receptor agonist isoproterenol, induced the recruitment of RAB18 to the surface of LDs in 3T3-L1 cells, where silencing of RAB18 impaired insulin-stimulated lipogenesis and its overexpression enhanced lipogenesis (Pulido et al. 2011). Together, these data suggest that RAB18 plays a role in both lipolysis and lipogenesis in 3T3-L1 adipocytes. In order to better understand the role of RAB18 in the regulation of lipid metabolism in primary adipocytes, in the current study, we evaluated the effects of age, fat location, metabolic status, and hormonal milieu on RAB18 expression (mRNA and protein) in rodent white adipose tissue (WAT).

Materials and methods

Reagents

Rabbit anti-rat RAB18 was purchased from Calbiochem (Barcelona, Spain). Anti-mouse adiponectin was from Chemicon Int. (Temecula, CA, USA) and rabbit anti-human CD45 was from Santa Cruz Biotechnology. TRIzol Reagent and DMEM were from Invitrogen Corp., PowerScriptTM reverse transcriptase from Clontech, SYBR Green-tag DNA polimerase from Bio-Rad, dNTPs from Ecogen (Madrid, Spain), and random primers from GE Healthcare (Buckinghamshire, UK). Normal goat serum and Envision Method Kit were from Dako (Glostrup, Denmark), ECL Plus Detection System from GE Healthcare, and anti-rabbit Alexa488-conjugated secondary antibody from Invitrogen Corp. Unless otherwise indicated, all other reagents were purchased from Sigma–Aldrich.

Animal studies

Sprague Dawley rats and GH-deficient dwarf and wild-type Lewis rats were obtained from Harlan Iberica (Barcelona, Spain). C57BL/6j, ob/ob mice and their lean littermate controls, as well as New Zealand obese (NZO) mice and their controls, were obtained from Jackson Laboratories (Bar Harbor, ME, USA). Experimental procedures were approved by the Institutional Animal Care and Use Committee. Animals had free access to standard rodent chow and tap water unless otherwise specified. Post mortem, subcutaneous (obtained from the inguinal fat pad), and visceral (obtained from the peri-gonadal fat pad) WAT samples were either processed for immuno-histochemistry and/or frozen in liquid nitrogen and stored at −80°C until analysis of Rab18 mRNA levels by RT-PCR or RAB18 protein content by immunoblotting, as described below.
**Ad libitum-fed Sprague Dawley rats** At postnatal (P) day 15, 45, 60, or 90, male and female rats were killed under fed conditions and WAT samples were excised and processed for immunohistochemistry or frozen for protein or mRNA analysis, as described below. In addition, in order to investigate the presence of RAB18 in the two components of adipose tissue, mature adipocytes and the stromal-vascular fraction (SVF; Frayn et al. 2003) were freshly isolated from WAT of P90 male Sprague Dawley rats, as described previously (Peinado et al. 2010). After separation, the two fractions were frozen in liquid nitrogen and stored at −80°C for analysis of Rab18 mRNA or protein content or freshly isolated mature adipocytes were processed for immunohistochemistry, as described below.

**Fasted mice** Visceral and subcutaneous WAT was collected from 10-week-old, C57bl/6J mice under ad libitum-fed conditions or after a 12-h fast (food removed at 1900 h) or 24 and 48 h (food removed at 0700 h). Samples were stored at −80°C until analysis of RAB18 mRNA and protein.

**Diet-induced and genetically obese mice** WAT was collected from C57Bl/6J mice maintained on a high-fat (fat = 60 kcal%, carbohydrate = 20 kcal%, protein = 20 kcal%; D12492, Research Diets, Brunswick, NJ, USA) or a low-fat diet (fat = 10 kcal%, carbohydrate = 70 kcal%, protein = 20 kcal%; D12450B, Research Diets) for 10 or 16 weeks (diet starting at 4 weeks of age), as previously reported (Luque & Kineman 2006). In addition, WAT samples from a model of monogenic obesity, ob/ob mice (10 weeks of age), or from a model of polygenic obesity, NZO mice (10 weeks of age), were also examined. We also analyzed the effect of 48-h fasting in NZO mice. Samples were stored at −80°C until analysis of Rab18 mRNA and protein.

**Hypophysectomized rats** Male Sprague Dawley rats were either sham-operated or hypophysectomized and WAT was collected 4 weeks after surgery as described previously (Barreiro et al. 2002).

**Growth hormone deficient dwarf rats** WAT was collected from 7-week-old controls and GH-deficient Lewis male rats, commonly referred to as spontaneous dwarf (SDR). In these rats, circulating GH levels are undetectable, due to a spontaneous point mutation in the GH gene resulting in a premature stop codon (Charlton et al. 1988).

**Adrenalectomized rats** Sprague Dawley rats were either sham-operated or bilaterally adrenalectomized, as previously reported (Barreiro et al. 2002). Both sham-operated and adrenalectomized rats received a daily s.c. administration of 40 µg dexamethasone dissolved in 200 µl saline or 200 µl saline alone for 8 days. Thereafter, sampling of WAT was carried out.

**Hypo- and hyper-thyroid rats** Hypothyroidism was induced in male Sprague Dawley rats by adding 0.1% 3-aminotriazole to the drinking water, and hyperthyroidism was induced by chronic s.c. administration of 100 µg/day l-thyroxine sodium salt pentahydrate, as previously reported (Lopez et al. 2001). The efficiency of the treatments was confirmed by evaluating the circulating levels of T4 and T3. Data from l-thyroxine-treated animals has been recently published (Varela et al. 2012). Administration of amino-triazole significantly decreased plasma levels of both T4 (107 ± 9 vs 56 ± 7 nmol/l in control animals and treated animals respectively; P<0.01) and T3 (1.63 ± 0.1 vs 0.56 ± 0.09 nmol/l in control animals and treated animals respectively; P<0.001).

**Immunohistochemistry**

Immunohistochemical detection of Rab18 was carried out in paraffin-embedded sections of visceral fat from adult (P90) male Sprague Dawley rats, which were processed as described previously (Archanco et al. 2007). Briefly, tissue sections were pre-incubated (1 h at room temperature) with 1:20 normal goat serum in 0.05 M Tris–HCl buffer, 0.5 M NaCl, pH 7.6 (TBS), and exposed overnight (4°C) to anti-rat RAB18 antibody (1:500). The signal was detected using the Envision Method Kit consisting of a goat anti-rabbit IgG secondary antibody coupled to a peroxidase-labeled dextran polymer. Peroxidase activity was revealed using 0.03% 3,3-diaminobenzidine. The reaction was stopped by a wash with TBS. Finally, sections were dehydrated and mounted with a mixture of distyrene, a plasticizer, and xylene (DPX). Negative control slides without primary antibody were included to assess nonspecific staining.

Immunohistochemical detection of Rab18 was also performed in freshly isolated mature adipocytes fixed with 4% paraformaldehyde (15 min) and then incubated with PBS containing 0.1% saponin and 1%...
BSA (1 h at RT). Cells were exposed overnight to anti-rat RAB18 antibody (1:500), followed by a 2-h incubation with anti-rabbit Alexa594-conjugated secondary antibody (1:500). The signal was visualized using a TCS-SP2-AOBS confocal laser scanning microscope (Leica Corp., Heidelberg, Germany).

**Western blot analysis**

Samples of total adipose tissue, mature adipocytes, and SVF were disrupted in lysis buffer (20 mM Tris, pH 7.4, 150 mM NaCl, 1% Triton, and protease inhibitors). Total protein (150 μg/lane) was loaded in 12.5% SDS–PAGE gels and transferred to nitrocellulose membranes. After Ponceau staining to ensure equal sample loading, membranes were blocked for 1 h with 5% dried milk in TTBS (TBS buffer with 0.05% Tween 20). Thereafter, membranes were incubated with primary antibodies (rabbit anti-rat RAB18, rabbit anti-mouse adiponectin, or rabbit anti-human CD45) overnight at 4°C, washed and incubated with peroxidase-conjugated secondary antibodies for 1 h at room temperature. Rabbit polyclonal anti-β-actin antibody was employed as a loading control. The signal was detected using the ECL Plus Detection System (GE Healthcare).

**Real-time RT-PCR**

Total RNA was isolated from WAT samples using TRIzol reagent. To assess RNA quality, the 260 nm/280 nm absorbance ratio was measured and samples were only used when this ratio was ~2. RT reactions were carried out in a 20 μl final volume by adding 2 μg total RNA, 1 μl PowerScript, 10 μM dNTP mixture, and 250 ng random hexamer primers. The reaction mixtures were incubated at 70°C for 10 min and at 42°C for 1 h and then heated at 70°C for 15 min and cooled on ice. Real-time RT-PCR was performed using SYBR Green tagging quantification in an iCycler IQ PCR detection System (Bio-Rad) according to the manufacturer’s instructions. The specific primers used were 5′-CTCTGAAGATCCTCATTGG-3′ sense and 5′-CTCTGAAGATCCTCATTGG-3′ antisense for rat Rab18 and 5′-CTCTGAAGATCCTCATTGG-3′ sense and 5′-CTCTGAAGATCCTCATTGG-3′ antisense for mouse Rab18. Both primer pairs amplify 185 bp fragments. As endogenous reference genes, the hypoxanthine phosphoribosyltransferase gene (Hprt) (for studies in rat adipose tissue) or the 18S small subunit ribosomal RNA gene (for studies in mouse adipose tissue) were amplified in parallel to Rab18 using the specific primers 5′-AGTCCCCAGCGTGATTAG-3′ sense and 5′-CTCTGAAGATCCTCATTGG-3′ antisense, which amplify a 139 bp fragment of the rat Hprt gene, and 5′-CCCATCTGACGCTGATTAG-3′ sense and 5′-CTCTGAAGATCCTCATTGG-3′ antisense, which amplify a 137 bp fragment of the mouse 18S rRNA gene. PCRs consisted of an initial activation and denaturing cycle at 95°C for 5 min, followed by 35 cycles at 95°C/30 s, 60°C/30 s, and 72°C/30 s. The amount of PCR products formed in each cycle was estimated on the basis of SYBR Green I fluorescence dye. Controls consisting of reaction mixture without cDNA were negative in all runs. cDNAs were sequenced to ensure that the correct mRNA transcripts were quantified (Central Sequencing Service, University of Cordoba, Cordoba, Spain).

Calculation of relative expression levels of the different transcripts was performed based on the cycle threshold (Ct) method. Thus, the Ct value for each sample was calculated using the iCycler iQ real-time PCR detection system software with an automatic fluorescence threshold setting. Reactions were performed in triplicate. Standard curves were constructed for Rab18 and the housekeeping genes and amplification profiles were analyzed off-line using the iCycler IQ optical system software 3.1 (Bio-Rad).

**Statistical analysis**

The number of animals used in each experimental setting is indicated in the figures. In each experiment, all samples were processed at the same time in order to reduce inter-experimental variability. Single comparisons were performed using Student’s t-test. Multiple comparisons were assessed by one-way ANOVA followed by a Newman–Keuls test for parametric data or by Kruskal–Wallis followed by a Dunn’s test for nonparametric data using GraphPad Prism 4 (La Jolla, CA, USA). Differences were considered statistically significant if P<0.05.

**Results**

**Distribution of Rab18 in adipose tissue**

Immunohistochemical examination of WAT sections or isolated mature adipocytes revealed that Rab18 immunoreactivity was present at the surface of the characteristic large unilocular LD (Fig. 1A and B). Rab18 immunoreactivity also showed a diffuse distribution within the thin layer of cytoplasm between the LD and the cell surface in isolated adipocytes (Fig. 1B).
Immunoblotting confirmed the presence of Rab18 immunoreactivity in mature adipocytes isolated from visceral and subcutaneous WAT, which also showed a strong immunosignal for the adipocyte marker, adiponectin, but not for the leukocyte marker CD45 (Fig. 1C). Rab18 immunosignal was also noticeable in the SVF of subcutaneous and visceral fat, characterized by the expression of CD45 (Fig. 1C).

**Influence of age and gender on Rab18 expression**

In WAT from both male and female rats, Rab18 expression was undetectable at P15 but was clearly detectable at P45, where levels remained constant up to P90, with no apparent sex differences (Fig. 2).

**Effect of fasting on WAT Rab18 mRNA expression**

As shown in Fig. 3A, Rab18 transcripts in visceral WAT increased with fasting, in a time-dependent manner, reaching a maximal value after 48 h of fasting (13-fold increase compared with control animals). Rab18 expression levels in subcutaneous WAT also increased after 12 h of fasting (15-fold increased compared with ad libitum-fed mice). However, in contrast to that observed in visceral fat, Rab18 mRNA levels in subcutaneous WAT tended to return to fed levels with prolonged fasting (24 and 48 h; Fig. 3B). Similar results were observed when RAB18 protein content in samples from fasted mice was assessed by immunoblotting (Fig. 3C).

**Effects of diet-induced and genetic obesity on Rab18 expression**

Ten weeks of HFD increased visceral WAT Rab18 mRNA levels to sixfold that observed in LFD controls (Fig. 4A). Likewise, Rab18 mRNA levels in subcutaneous WAT were tenfold higher in mice fed HFD than in mice fed LFD (Fig. 4B). After 16 weeks of HFD, visceral and subcutaneous WAT Rab18 transcript levels remained elevated (Fig. 4C and D), although differences between HFD-fed and LFD-fed animals were less pronounced than in the 10-week group. Rab18 mRNA content in visceral and subcutaneous fat from obese, leptin-deficient mice (i.e. ob/ob mice) was also dramatically elevated in both visceral (Fig. 4E) and subcutaneous (Fig. 4F) WAT, compared with lean controls (400- and 300-fold respectively). The differences in Rab18 mRNA levels in visceral WAT were largely matched by changes in protein expression; however, RAB18 protein levels in subcutaneous WAT did not differ between LFD and HFD groups (Fig. 4G).
RAB18 protein content was higher in both visceral and subcutaneous adipose tissue of ob/ob animals than in their corresponding control counterparts (Fig. 4H). As shown in Fig. 4G and H, adiponectin protein expression in adipose tissue decreased in both diet-induced and genetic obesity, with the most pronounced differences observed in visceral WAT.

Similar to that found in ob/ob mice, visceral WAT samples from NZO mice also showed increased levels of Rab18 expression compared with their corresponding lean control animals (14-fold higher; Fig. 5A), although differences were less pronounced than those observed in ob/ob mice. Interestingly, similar to that observed in lean mice (Fig. 3A), 48-h fasting increased Rab18 mRNA content in visceral WAT of NZO mice (Fig. 5B).

**Hormonal regulation of WAT Rab18 gene expression**

Given the involvement of pituitary hormones, and those produced by other endocrine tissues controlled by this gland, in the regulation of adipocyte lipid metabolism (Schaffler et al. 2005, 2006), we explored the impact of disrupting pituitary hormone output on Rab18 mRNA levels in WAT. As illustrated in Fig. 6A, Rab18 expression was 1.9-fold higher in hypophysectomized rats than in sham-operated animals, thus indicating that pituitary hormones, either directly or via their actions on peripheral glands, could be involved in the regulation of Rab18 gene expression in WAT. To ascertain which pituitary hormone(s) are responsible for maintaining WAT Rab18 expression, Rab18 mRNA levels were measured in GH-deficient (dwarf; Fig. 6B), gonadectomized (Fig. 6C and D), adrenalectomized (with and without dexamethasone treatment; Fig. 6E), and hyperthyroid rats (Fig. 6F) and found not to differ from their respective controls. However, of note, hypothyroidism increased WAT Rab18 mRNA (Fig. 6F) and protein (Fig. 6G) levels compared with euthyroid rats.

**Discussion**

In this study, we demonstrate that Rab18, which is associated with the surface of LDs in primary adipocytes, is regulated in WAT in an age- and depot-dependent manner, in relation to the energy status (i.e. fasting and diet-induced or genetic obesity), and thyroid function. These observations, mostly based on mRNA data, are in agreement with previous functional results supporting the participation of Rab18 in the regulation of lipid metabolism in adipose tissue (Pulido et al. 2011).
During postnatal development, Rab18 expression was undetectable in visceral WAT from P15 rats, thereby suggesting that this GTPase is not required for the initial expansion of adipose tissue, which occurs in early postnatal life (Gregoire et al. 1998). However, Rab18 expression exhibited a marked increase during puberty, reaching the highest levels at P45 and remaining elevated thereafter. This age-related increase in Rab18 expression coincides with the period of highest adipocyte hypertrophy in rat WAT, which occurs between P40 and P80 and is characterized by an increase in the lipogenic activity and a decline in the lipolytic rate of adipocytes (Tsujikawa & Kimura 1980, Herrera & Amusquivar 2000). Accordingly, it has been reported that Rab18 mRNA expression and protein content increased in 3T3-L1 adipocytes as these cells accumulated lipids during differentiation (Pulido et al. 2011). This may account for the presence of Rab18 immunoreactivity in the SVF of both subcutaneous and visceral fat, as this component of WAT contains a population of preadipocytes (Frayn et al. 2003). Nevertheless, we cannot exclude the possibility that other cell types in the SVF can also contribute to the changes in Rab18 mRNA content observed in this study. It is notable that in spite of the sexual differences in the lipolytic and lipogenic rates of adipose tissue, which have been largely attributed to the influence of sex steroids (Shi & Clegg 2009, Shi et al. 2009), no gender-related differences in Rab18 mRNA expression were observed at any of the ages tested, even in periods of increased sex steroids levels (i.e. puberty). In line with this, we observed that gonadectomy did not modify Rab18 transcript levels in WAT.

Our study also showed that fasting induced a strong, time-dependent increase in Rab18 gene expression in WAT, which supports the participation of this protein in the adaptive response of adipose tissue to maintain whole-body homeostasis. During fasting, increased concentrations of circulating catecholamines and glucocorticoids promote mobilization of lipid stores from adipose tissue (Langin 2006, Rose et al. 2010). Glucocorticoids do not appear to play a major role in the regulation of Rab18 expression in WAT as much as administration of dexamethasone to either adrenalectomized or sham-operated animals did not alter the expression levels of this GTPase. Additionally, exposure of 3T3-L1 adipocytes to dexamethasone had no effect on Rab18 mRNA levels (Pulido et al. 2011). On the other hand, the β-adrenergic agonist isoproterenol, besides increasing the association of this protein to the LD surface, enhanced Rab18 transcript and protein content in 3T3-L1 cells (Pulido et al. 2011). Taken together, these data suggest the involvement of Rab18 in β-adrenergic receptor-stimulated lipolysis during short-term energy deficit. Our results on the distinct time-course profile of Rab18 mRNA expression exhibited by subcutaneous and visceral fat in response to fasting also support this notion. To be more specific, whereas Rab18 mRNA content in visceral fat increased steadily over a 48-h fasting period, subcutaneous fat exhibited a sharp, but transient increase in Rab18 mRNA levels in response.
to food deprivation. Moreover, changes in RAB18 protein content in adipose tissue samples induced by fasting were similar to those observed for Rab18 mRNA. These observations are in accordance with the higher expression levels of β-adrenergic receptors and increased β-adrenergic stimulation of lipolysis exhibited by visceral, compared with subcutaneous, WAT (Hoffstedt et al. 1997, Umekawa et al. 1997, Portillo et al. 2000). Moreover, visceral fat is more responsive to fasting conditions than subcutaneous fat with regard to the induction of the lipolytic enzymes, hormone-sensitive lipase (HSL; Akesson et al. 2003), and adipose triglyceride lipase (ATGL; Palou et al. 2010).

In this context, the results of the current study raise the possibility that Rab18 induction may help to sustain the higher lipolytic activity of visceral vs subcutaneous fat in response to energy deprivation.

Remarkably, we observed that Rab18 mRNA levels in WAT were strongly increased in obesity induced by HFD, hyperphagia due to leptin deficiency (ob/ob mice), or to polygenic alterations (New Zealand mice). A similar trend was observed for Rab18 protein expression in WAT from HFD mice and ob/ob mice. In agreement with these findings, we have recently shown that obese humans also exhibit enhanced Rab18 expression levels in both visceral and subcutaneous fat (Pulido et al. 2011). Together, these data suggest that Rab18 may be involved in the metabolic changes occurring in fat under conditions of energy excess. In particular, HFD-induced obesity is characterized by increased basal catecholamine-induced lipolysis (Jocken & Blaak 2008, Gaidhu et al. 2010). Likewise, adipose tissue of ob/ob mice exhibits higher rates of net lipolysis than their lean controls (Turner et al. 2007). Intriguingly, although gene...
expression studies have commonly reported reduced or normal expression levels of lipogenic genes in ob/ob mice (Nadler et al. 2000), measurement of triacylglyceride synthesis and de novo lipogenesis has revealed that adipose tissue capacity for lipid anabolism and storage is markedly increased in these animals (Turner et al. 2007). Notably, overexpression of Rab18 stimulates both lipolytic and lipogenic rates in 3T3-L1 adipocytes (Pulido et al. 2011). According to these data, it is tempting to propose that this protein could participate in the regulation of lipid mobilization from (i.e. lipolysis) and to (i.e. lipogenesis) LDs in response to increased adiposity.

Besides the marked effects on Rab18 mRNA levels triggered by changes in the energy status, disruption of pituitary signaling also altered the expression of this GTPase in WAT. Specifically, WAT Rab18 mRNA levels increased significantly in hypophysectomized rats, indicating that pituitary hormones, either directly or through their action on target glands, regulate the expression of this GTPase. However, most of the experimental procedures aimed at disrupting different pituitary-related endocrine axes were ineffective at altering Rab18 expression, with the exception of hypothyroidism. Specifically, hypothyroidism upregulated Rab18 mRNA expression, as hypophysectomy did, and also increased WAT Rab18 protein content. Interestingly, both hypophysectomy and hypothyroidism have been shown to decrease basal and catecholamine-stimulated lipolysis (Malbon et al. 1978, Mills et al. 1986, Yang et al. 1996), where lipogenesis is unchanged or increased (Correze et al. 1982, Baht & Saggerson 1988, Hausman et al. 1999). In all, our data suggest that hypophysectomy-induced effects on Rab18 mRNA expression may be mediated by the decrease in circulating thyroid hormones resulting from the lack of pituitary TSH. Nonetheless, we cannot exclude the possibility that other pituitary hormones (or lack thereof) not investigated in this study and known to act on adipose tissue, such as PRL (Brandebourg et al. 2007), might also be responsible, at least in part, for the effects observed in hypophysectomized animals.

From the findings reported herein, it is worthy to highlight that Rab18 expression is upregulated in rodent WAT from both visceral and subcutaneous depots in response to different metabolic situations, which might represent an adaptive response to overcome the alterations in lipid metabolism occurring under such conditions. Thus, Rab18 is induced in states of augmented lipogenesis and decreased lipolysis, as occurs during postnatal development, when the pituitary/thyroid axis is disrupted, in states of increased lipolytic activity mediated by sympathetic signaling triggered by fasting, and in states of increased lipogenic and lipolytic activities as a result of obesity. These observations are largely in accordance with our previous functional data in 3T3-L1 adipocytes demonstrating the participation of Rab18 as a common intracellular mediator of lipolysis and lipogenesis (Pulido et al. 2011). Together, these data suggest the involvement of Rab18 in the regulation of lipid metabolism in adipose tissue under physiological and pathological conditions and provide further support to the idea that this may occur by controlling both the lipolytic and the lipogenic activities of adipocytes.


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