Regulation of adipose tissue leptin secretion by α-melanocyte-stimulating hormone and agouti-related protein: further evidence of an interaction between leptin and the melanocortin signalling system

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Abstract

The central role of the melanocortin system in the regulation of energy balance has been studied in great detail. However, the functions of circulating melanocortins and the roles of their peripheral receptors remain to be elucidated. There is increasing evidence of a peripheral action of melanocortins in the regulation of leptin production by adipocytes. Here we investigate the interaction of α-melanocyte stimulating hormone (α-MSH) and agouti-related protein (AgRP) in the regulation of leptin secretion from cultured rat adipocytes and examine the changes in circulating α-MSH and AgRP in lean and obese rodents after hormonal and energetic challenge. Leptin secretion (measured by ELISA) and gene expression (by real-time quantitative PCR) of differentiated rat adipocytes cultured in vitro were inhibited by the administration of α-MSH (EC50=0·24 nM), and this effect was antagonised by antagonists of the melanocortin receptors MC4R and MC3R (AgRP and SHU9119). The presence of MC4R in rat adipocytes (RT-PCR and restriction digest) supports the involvement of this receptor subtype in this interaction. Leptin administered to ob/ob mice in turn increases the release of α-MSH into the circulation, suggesting a possible feedback loop between the site of α-MSH release and the release of leptin from the adipose tissue. However, the physiological significance of this putative feedback probably depends upon the underlying state of energy balance, since in the fasting state low plasma α-MSH is paralleled by low plasma leptin.


Introduction

Leptin secreted by adipose tissue regulates energy balance principally through its hypothalamic receptors. Although leptin receptors in the hypothalamus are known to interact with pathways involving a number of neuropeptides, neurons originating in the arcuate nucleus that release α-melanocyte stimulating hormone (α-MSH) in the paraventricular nucleus form an important part of this system. The melanocortin receptors (MC-Rs) thought to be involved in the energy balance signalling are the MC4R and MC3R. An integral part of this system appears to be another leptin-sensitive pathway releasing agouti-related protein (AgRP), an MC4/3R antagonist. Leptin thus stimulates α-MSH and inhibits AgRP in a coordinated manner to regulate food intake. For this reason much attention has been focused on the role of the melanocortin system in obesity (see Ahima et al. 2000, MacNeil et al. 2002, Zimanyi & Pelleymounter 2003).

Melanocortin peptides are generated from a common precursor glycoprotein, pro-opiomelanocortin (POMC), by post-translational processing. The POMC gene is expressed at high levels in the arcuate nucleus of the hypothalamus. It is also expressed in the anterior and intermediate lobes of the pituitary, and at lower levels in a wide variety of mammalian peripheral tissues. MSHs are now recognised to have a variety of physiological functions, reflecting the wide distribution of their
corresponding MC-Rs (Zimanyi & Pelleymounter 2003). Five subtypes of MC-Rs have been identified (MC1–5R) with discrete pharmacological properties and tissue distribution, including several subtypes present in adipose tissue (MC1R, MC2R, MC4R and MC5R) (Boston & Cone 1996, Chagnon et al. 1997). Two endogenous antagonists of MC-Rs have been identified, agouti and AgRP (Zimanyi & Pelleymounter 2003). AgRP is also present in the systemic circulation (Li et al. 2000). The function of circulating AgRP along with the peripheral roles of melanocortins and MC-Rs remain to be elucidated. α-MSH has been shown to interact with several peripheral organ systems. It is a potent lipolytic agent in a number of species. This was first identified over 25 years ago in rabbits (Kastin et al. 1975), and recently similarly potent lipolytic activity has been shown in rodents, providing support for a role of α-MSH in the integration of peripheral metabolism with central appetite regulation (Forbes et al. 2001). α-MSH has been reported to regulate insulin secretion, inhibit the effects of cytokines and apoptosis, stimulate corticosterone release, and to have a role in thermogenesis (Vinson et al. 1983, Shimizu et al. 1995, Rajora et al. 1997, Forbes et al. 2001, Jo et al. 2001). In addition there is increasing evidence for a peripheral action of melanocortins in the regulation of leptin production by adipocytes (Hoggard et al. 2001, Norman et al. 2003).

In this paper we investigate the interaction of α-MSH and AgRP in the regulation of leptin secretion from cultured rat adipocytes and examine the changes in circulating α-MSH and AgRP in relation to energy balance and leptin signalling in both lean and obese rodents. This study, part of which has been published in abstract form (Hoggard et al. 2001), extends the recent reports of an interaction between leptin and melanocortins in murine adipocyte culture (Norman et al. 2003).

Materials and methods

Animals

Lean (+/?) and obese (ob/ob) Aston mice (11 weeks), Hooded Lister rats (14 days), and lean (+/?) and obese Zucker (fa/fa) rats (14 weeks) were all drawn from colonies maintained at the Rowett Research Institute. Food (Biosure; Special Diets Services, Witham, Essex, UK) and water were available freely unless stated. All animals were killed by cervical dislocation in the middle of the light phase. To examine the effects of food deprivation, lean (+/?) Aston mice were fasted for 24 h. To test the effect of leptin challenge, obese (ob/ob) Aston mice were injected i.p. with either leptin (1·7 mg/kg body weight) or saline. Animals were killed by cervical dislocation 1 h after injection. Tissues were dissected and frozen immediately in liquid nitrogen prior to transfer to −80°C where they were stored until extraction of RNA. All procedures were licensed under the UK Animals (Scientific Procedures) Act of 1986 and received ethical approval from the Rowett Research Institute’s Ethical Review Committee.

Hormones

α-MSH (Sigma; M4135), [Nle⁴,DPhe⁷]-αMSH (Sigma; M8764) and SHU9119 (Sigma; M4603) were purchased from Sigma-Aldrich Company Ltd, Poole, Dorset, UK. AgRP (005–57) was purchased from Phoenix Peptides, Belmont, CA, USA.

Cell culture

Fibroblastic preadipocytes were isolated from adipose tissue as previously described (Mitchell et al. 1997). The inguinal fat pads from 14-day-old male Hooded Lister rats (eight per group) were removed under sterile conditions. The resultant cell preparation (mainly fibroblastic preadipocytes) was adjusted to a density of 1·5×10⁵ cells/ml in Medium 199 with 10% fetal calf serum (Gibco BRL); 1·5 ml volumes were plated onto six-well plates. After 4 days in culture at 37°C in an atmosphere of 5% CO₂, differentiation was induced by the addition of medium supplemented with isobutylmethylxanthine (0·5 mM; Sigma), dexamethasone (0·25 mM; Sigma) and insulin (10 mg/ml; CP Pharmaceuticals, Wrexham, UK). After 48 h, the induction medium was removed and replaced by Medium 199 containing 10% fetal calf serum supplemented with insulin (10 mg/ml) alone. This medium was changed every 2 days. Eight days post-differentiation, cells were incubated with or without test hormone. Following incubation, samples were retained and centrifuged at
150 g for 10 min; the supernatant was stored at −80 °C for the assay of leptin. Cells were lysed directly on the plate for determination of protein by the Bio-Rad Protein Assay (Cat. no. 500–0006) or for RNA extraction (RNAqueous-4 PCR isolation kit; Ambion Inc.), prior to leptin mRNA determination.

**Leptin ELISA**

Plasma leptin was determined by an in-house chemiluminescence ELISA, as previously described (Hardie et al. 1996, Crabtree et al. 2000). Chemiluminescence was achieved by the addition of 100 µl/well CDP-Star substrate in enhancer solution (Sapphire-II enhancer in DEA buffer; 1:10; Applied Biosystems, Warrington, UK) and quantified in an MLX luminometer (Dynex, Worthing, West Sussex, UK) in terms of relative light units. Recombinant murine leptin standards (NIBSC, South Mimms, Hertfordshire, UK) were run in triplicate and appropriate blanks of culture medium were included. Results were expressed as a percentage of the control after correction for minor differences in protein levels in the wells. Each incubation was carried out in six individual wells.

**α-MSH detection**

Plasma α-MSH was detected using a commercial kit according to the manufacturer’s instructions (Euro-Diagnostica IDS Ltd, Boldon, Tyne and Wear, UK). All samples were run in duplicate. The minimum level of detection of α-MSH was 3 pM and the intra- and inter-assay coefficients of variation of the assay were 11·8 and 13·0% respectively. The cross-reactivity with other POMC peptides (adrenocorticotrophic hormone [ACTH] (1–24), ACTH (1–39), β-MSH and γ-MSH) was stated to be <0·002%.

**AgRP detection**

Plasma AgRP was detected using a commercial kit according to the manufacturer’s instructions (Phoenix Peptide). All samples were run in duplicate. The detection range for AgRP was 1–128 pg/ml. Cross-reactivity with AgRP(83–131)-NH₂ (mouse), AgRP(83–132)-NH₂ (human) and AgRP form C was 100%. Cross-reactivity with orexin A, orexin B (human), orexin B (rat, mouse), leptin (human), leptin (mouse), α-MSH, glucagon-like peptide (GLP)-1(7–37), GLP-2 and neuropeptide Y (human) was 0%.

**RT-PCR of MC4R and MC5R**

RNA was extracted using an RNAqueous-4 PCR isolation kit for DNA-free RNA, following the manufacturer’s instructions (AMS Biotechnologies, Abingdon, Oxon, UK). Reverse transcription was carried out on 5 µg RNA using the Superscript Preamplification System (Gibco BRL, Maidenhead, Berkshire, UK) according to the manufacturer’s instructions. The cDNA was amplified by PCR using rodent-specific primers for the MC-R. The quality of each cDNA and mock cDNA was determined by the relative level of amplification of the rodent β-actin gene (542 bp product) as previously described (Hoggard et al. 1997). The primers were as follows: MC4R and MC5R were as previously described (Langouche et al. 2001); MC4R (GenBank Accession number U67863), forward 5’-TTTCATCTTAGTCTG GCT-3’ (nucleotides 389–408), reverse 5’T-GGCGCGATACTGT GCAAGCT-3’ (nucleotides 694–672), product size 305 bp; MC5R (GenBank Accession number L27081), forward 5’-TCTTCT TTGTGGGCAGCC TAG-3’ (nucleotides 469–487), reverse 5’T-CAGG GCGTAGAAGATGGTGATGTAC-3’ (nucleotides 696–672), product size 228 bp. All primers were synthesised by Sigma-Genosys Ltd, Cambridge, UK. PCR was performed on a Touchdown thermal cycler (Thermo Hybaid, Ashford, Middlesex, UK) using the following conditions for both sets of MC-R primers, 95 °C (10 min) 1 cycle, 95 °C (20 s), 60 °C (20 s), 72 °C (30 s) for 35 cycles followed by a final extension of 72 °C for 7 min. Agarose gel electrophoresis (2%) in the presence of ethidium bromide confirmed the presence of a single band of the expected size. To confirm the identity of the amplified products, restriction analysis was carried out on each PCR product using specific restriction enzymes as previously described (Langouche et al. 2001). SsP1 was specific for MC4R cDNA and MboI was specific for MC5R cDNA. All cDNA samples were amplified by β-actin primers (data not shown), demonstrating the viability of the RNA.
Real-time quantitative PCR (Taqman system)

RNA was extracted using an RNAqueous-4 PCR isolation kit for DNA-free RNA, following the manufacturer’s instructions (AMS Biotechnologies). RNA was accurately quantified on the Agilent 2100 Bioanalyser (Agilent Technologies, South Queensferry, West Lothian, UK). This also shows the quality of the RNA extracted from the tissues. Only RNA showing no degradation was processed further. Reverse transcription was carried out on 5 μg RNA using the Superscript Preamplification System (Gibco BRL) according to the manufacturer’s instructions. The cDNA was amplified by PCR using rodent-specific primers for the leptin gene. The primers were as follows: leptin, forward 74–97, 5′-TTGTCACCAGGATCAATGACATTT-3′, GenBank U48849; reverse 157–179, GACAACTCAGAATGGGGTGAAG-3′, GenBank U48849; and FAM-labelled Taqman probe, 99–121, ACACACGCAGTCCGTATCCGCCA, Genbank U48849.

The cDNAs were also amplified using rodent ribosomal 18S RNA and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) control primers (PE Applied Biosystems, Inc., Foster City, CA, USA), following the manufacturer’s instructions. PCR was performed on an Applied Biosystems ABI Prism 7700 sequence detection system under the following conditions; 50°C (2 min) for 1 cycle, 95°C (10 min) 1 cycle, 95°C (15 s), 60°C (1 min) for 40 cycles. All samples were run in triplicate and were all within the standard curve. Values were standardised to both 18S and GAPDH and expressed as a percentage of the control.

Statistics

Between-group comparisons were made using unpaired Student’s t-tests. Results were considered statistically significant if the P value was less than 0.05.

Results

Regulation of leptin secretion and expression in primary culture of differentiated rat adipocytes in vitro

The rat adipocytes were treated at day 8 post-differentiation. Twenty-four hours later the culture medium was harvested for the measurement of leptin, using a sensitive chemiluminescent ELISA and cells extracted for protein determination.

Effect of α-MSH on leptin secretion into the culture medium

Differentiated rat adipocytes were incubated for 24 h with five different concentrations of α-MSH (0.015, 0.15, 1.5, 15 and 150 nM) and compared with control cells (Fig. 1). Addition of α-MSH resulted in a dose-dependent decrease in leptin secretion into the culture medium. The EC₅₀ for the reduction of leptin secretion by α-MSH was determined to be approximately 0.24 nM (Fig. 1). This experiment was repeated several times with similar results.

Effect of a potent α-MSH analogue and α-MSH antagonists on leptin secretion into the culture medium

The effect of [Nle⁴, d-Phe⁷]-α-MSH, a potent analogue of α-MSH, on leptin secretion into the medium was determined on differentiated primary adipocytes (Fig. 2). Cells were incubated with [Nle⁴, d-Phe⁷]-α-MSH for 24 h and compared with control (medium only). [Nle⁴, d-Phe⁷]-α-MSH at 15 nM decreased leptin secretion into the culture medium compared with the control adipocytes (P<0.05). This decrease in leptin secretion by [Nle⁴, d-Phe⁷]-α-MSH was inhibited by the addition of SHU9119 (1 μM) or AgRP (200 nM). Pre-incubation with the antagonists prior to the addition of α-MSH had the same effect on leptin secretion (data not shown). The addition of AgRP...
(200 nM) in the absence of α-MSH had no effect on the secretion of leptin. Similar results were obtained in two further separate experiments.

**Figure 2** Addition of the potent α-MSH analogue [Nle⁴, D-Phe⁷]-α-MSH (NP-MSH) at 15 nM to primary cultures of differentiated adipocytes for 24 h significantly (P<0.05) inhibits leptin protein secreted into the cell culture medium compared with untreated control cells, as determined by a leptin-specific ELISA. This inhibition of leptin by [Nle⁴, D-Phe⁷]-α-MSH is blocked by the addition of SHU9119 (1 µM) or AgRP (200 nM), which are both potent antagonists of α-MSH. AgRP (200 nM) treatment of the cells on its own has no effect on leptin secretion into the culture media. n=6 for each point, per cent of control±S.E.M. *P<0.05.

**Figure 3** The addition of α-MSH (1.5 or 15 nM) to differentiated primary cultures of rat adipocytes for 6 h significantly (*P<0.05) decreased leptin mRNA expression, when compared with the untreated control cells. Leptin mRNA was determined by real-time quantitative PCR using the Taqman assay and primers specific for leptin. Results are expressed as a percentage of the control, n=3 for each point±S.E.M, related to the expression of the house keeping gene 18S.

**Figure 4** RT-PCR ethidium bromide-stained gels showing the expression of MC4R (a and b) and MC5R (c) in primary cultures of rat differentiated adipocytes (P), rat adipose (A) tissue (epididymal), rat brain (Br), control PCR (Co), mouse brain, (Bm), MC4R band amplified from adipose tissue digested with SsP1 (D), and 100 bp ladder with the 300 bp fragment shown.

**Effect of α-MSH on leptin mRNA expression**

The addition of α-MSH (1.5 or 15 nM) to differentiated primary cultures of rat adipocytes for 6 h significantly (P<0.05) decreased leptin mRNA expression, as determined by the Taqman assay using primers specific for leptin mRNA (Fig. 3). α-MSH (1.5 nM) decreased leptin mRNA by approximately 40% when compared with the control cells (medium only).

**MC-R subtypes present in differentiated primary cultures of rat adipocytes and rat adipose tissue**

To determine if MC4R receptors are present in the primary cultures of rat adipocytes and in rat adipose tissue we used RT-PCR and primers specific to MC4R (Fig. 4a and b) or to MC5R as a control (Fig. 4c). Rat and mouse brain were used as control tissue for MC4R. Half of the MC4R PCR reaction was analysed on an ethidium bromide-stained agarose gel and the other half was digested with SsP1 restriction enzyme, which yields two PCR fragments of 119 and 186 bp, to confirm the identity of the product (Fig. 4b). Analysis shows that MC4R and MC5R are present in rat adipose tissue and in differentiated primary cultures of rat adipocytes.

Specificity of MC-R mRNA amplification was demonstrated. No β-actin band was amplified from the mock cDNA, indicating that the DNAse
treatment in the RNA extraction was complete and that any bands amplified from the RT-RNA were not as a result of contaminating DNA present in the preparation.

Circulating levels of α-MSH and AgRP in ob/ob mice and fa/fa rats

To investigate whether leptin regulates the release of α-MSH or AgRP into the circulation we examined plasma levels of α-MSH and AgRP in leptin-deficient ob/ob mice and leptin receptor-defective fa/fa rats, comparing the levels of these hormones with their lean littermates (Fig. 5A). Circulating α-MSH was significantly \((P<0.05)\) lower in leptin-deficient ob/ob mice, and leptin receptor-defective fa/fa rats \((P<0.05)\) compared with their respective lean littermates \((+/?)\). The administration of leptin i.p. at 1.7 mg/kg body weight to ob/ob mice increases circulating α-MSH compared with saline-treated controls after 1 h \((P<0.05)\) (Fig. 5A). No significant changes in plasma AgRP were observed in either ob/ob mice or fa/fa rats compared with their respective lean littermates (B). \(n=6\pm \text{S.E.M.}\)

Circulating levels of α-MSH and AgRP in 24 h food-deprived mice

To investigate whether α-MSH or AgRP have a role in the regulation of food intake, plasma levels of these hormones were determined in 24 h food-deprived mice (Fig. 6). Plasma α-MSH levels were significantly lower \((P<0.05)\) compared with the freely fed controls. There was no significant change in circulating AgRP, although there was a trend for AgRP levels to be higher in the food-deprived group.

Discussion

We have shown that expression of leptin, both mRNA and protein, is inhibited \textit{in vitro} by the
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administration of α-MSH to primary cultures of rat adipocytes, and that this effect is antagonised by MC4/3R antagonists. This inhibition of leptin by α-MSH in rat adipocytes (EC$_{50}$ = 0·24 nM) is much more pronounced that that previously reported in murine 3T3-L1 cells (EC$_{50}$ = 36 nM). This may be due to the presence of the MC4R in rat adipocytes, a receptor subtype that is also present in human adipose tissue, but which is absent in 3T3-L1 cells. Leptin administered to ob/ob mice in turn increases the release of α-MSH into the circulation, suggesting a possible feedback loop in vivo whereby increased leptin secretion from adipose tissue stimulates an increase in plasma α-MSH, which in turn inhibits leptin release. However, we propose that the physiological role of this putative homeostatic feedback varies with the state of energy balance, since in the fasting state low plasma α-MSH and low plasma leptin are co-incident.

At this stage it is not possible to say whether this inhibition of leptin expression is a direct effect on gene transcription or the result of the potent lipolytic action of α-MSH on adipose tissue (Forbes et al. 2001), as a reduction in cell size has been shown to be an important regulator of leptin expression from adipose tissue (Van Harmelen et al. 1998). However, in support of a direct effect, MC-Rs are coupled to adenylyl cyclases, and cAMP has been shown to strongly down regulate leptin mRNA expression in rat adipocytes in culture (Slieker et al. 1996).

Signalling in the POMC system involves five separate receptors, which respond to the different peptides in the system, but with a measure of cross-talk between them (Wikberg et al. 2000). A number of MC-Rs have been identified in adipose tissue, but there appear to be clear differences between species. In human adipose tissue, MC1R, MC2R, MC4R and MC5R have been shown to be expressed (Chagnon et al. 1997), which contrasts with murine adipose tissue where only the MC2R and MC5R have been reported (Boston & Cone 1996). α-MSH is unlikely to signal through MC2R, which is the ACTH receptor (Wikberg et al. 2000). Consequently, it has previously been suggested that in 3T3-L1 murine adipocytes α-MSH, and in addition ACTH, down regulate leptin expression via the MC5R (Norman et al. 2003). The presence of MC4R in rat adipocytes, but not murine 3T3-L1 cells, may contribute to the much lower EC$_{50}$ value of 0·24 nM reported here for the action of α-MSH on rat primary cultures of adipocytes compared with 36 nM for the action of α-MSH on leptin in 3T3-L1 cells. In support of α-MSH acting through the MC4R in rat adipocytes, both AgRP (200 nM) and SHU9119 (1 µM) block the inhibition of leptin secretion by α-MSH, consistent with their potent antagonism of α-MSH action at the MC3R and MC4R (Wikberg et al. 2000). The presence of the MC4R in rat adipose tissue may mean that this is a better model of human adipose tissue than mouse cells, since human adipocytes also express this receptor. Regardless of the receptor that α-MSH is signalling through, it clearly has an important signalling role in the regulation of leptin expression, at least in rodents and probably in humans too.

Agouti (100 nM), which acts in a similar manner to AgRP, has been shown to increase (approximately 1·6-fold) both leptin secretion and mRNA expression in differentiated 3T3-L1 murine adipocytes (Mynnatt & Stephens 2001). We therefore examined whether AgRP had a direct effect on leptin expression in primary culture of rat adipocytes. The addition of AgRP at supraphysiological levels to our primary culture of rat adipocytes (Mynatt & Stephens 2001). We therefore investigated whether leptin regulates the release of α-MSH or AgRP into the circulation. Consistent with this proposal, plasma α-MSH was decreased in both the leptin-deficient ob/ob mouse and the leptin receptor-defective fa/fa rat compared with their respective lean littermates. Further, administration of leptin to ob/ob mice stimulated an increase in circulating plasma α-MSH. Previous studies have shown similar effects (Forbes et al. 2001). This suggests some form of feedback loop between the peripheral secretion of leptin from adipose tissue and the release of α-MSH into the plasma. In keeping with this, POMC gene expression is decreased in the hypothalamus of the ob/ob mouse (Thornton et al. 1997). This is consistent with the hypothalamus being the source of plasma α-MSH. In contrast we observed no change in plasma AgRP in either the ob/ob mouse or the fa/fa rat compared with their respective lean littermates, suggesting that the release of AgRP into the circulation is not regulated by plasma leptin. This is in contrast to the
expression of AgRP in the arcuate nucleus of the hypothalamus, which has been shown to be regulated by leptin, suggesting that the hypothalamus may not be the source of circulating AgRP (Shutter et al. 1997). In humans and rodents the most abundant site of AgRP mRNA expression is the hypothalamus, followed by the adrenal glands (cortex and medulla). AgRP expression has also been reported in lung, kidney, testis, ovary and muscle tissues (Shutter et al. 1997, Takeuchi et al. 2000).

To investigate this putative feedback loop further we looked at the plasma levels of both α-MSH and AgRP in 24 h food-deprived mice. It would appear from these studies that the physiological significance of this feedback loop probably depends upon the underlying state of energy balance, since in the fasting state low plasma α-MSH is paralleled by low plasma leptin. A possible explanation is that in fasting the sympathetic system, which we have previously shown to be a key inhibitor of leptin (Trayhurn et al. 1998), overrides the α-MSH signal, maintaining a low plasma leptin level, which in turn results in a low plasma α-MSH. In addition the α-MSH signal may be further inhibited by the increase in AgRP observed in the fasting state (Li et al. 2000); although there was no significant change in circulating AgRP in response to a 24 h fast, the trend apparent in our data was in agreement with a previous report in 48 h food-deprived mice (Li et al. 2000). The physiological role of AgRP on adipose tissue MC4R may be questioned as the concentrations of AgRP shown in this study and in previous work are less than the \( K_i \) (2.5 nM) for AgRP at this receptor (Yang et al. 1999). However, circulating levels in vivo may not correlate with in vitro estimations of concentrations necessary to inhibit these receptors, particularly in view of the postulated role of syndecan-3 to bind and potentiate the effects of AgRP at MC-Rs in the hypothalamus (Reizes et al. 2001).

It should, however, be noted that the plasma levels of α-MSH reported here in the mouse, although within the physiological range for the regulation of lipolysis in vitro (Kastin et al. 1975), are lower than the previously reported EC\(_{50}\) value for the induction of leptin in murine 3T3 cells (Norman et al. 2003). The inhibition of leptin by basal levels of circulating α-MSH under physiological conditions in the mouse therefore requires further clarification.

Both plasma α-MSH (Katsuki et al. 2000) and AgRP (Katsuki et al. 2001) are increased in obese humans compared with lean subjects, suggesting that these circulating hormones may be involved in the regulation of food intake/body fat independently of any central effects, a hypothesis we have investigated further in rodents. We have shown that α-MSH regulates leptin expression in vitro, an effect which is antagonised by AgRP. This appears to form part of an integrative regulatory loop with central regulation of POMC gene expression, and subsequent release of α-MSH into the circulation. This putative feedback loop is a construct based on available evidence, but its regulatory capability in the normal, leptin-competent rodent under normal physiological conditions remains to be determined. In negative energy balance we suggest that the α-MSH signal is overridden by the sympathetic control of leptin production and lipolysis, reinforced by the antagonism of the α-MSH signal as a result of an increase in plasma AgRP.

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