High fat diet and body weight have different effects on cannabinoid CB₁ receptor expression in rat nodose ganglia

N.L. Clunya, E.D. Barbob, K. Mackiec, G. Burdyga, D. Richard, G.J. Dockray, K.A. Sharkey

Abstract

Energy balance is regulated, in part, by the orexigenic signaling pathways of the vagus nerve. Fasting-induced modifications in the expression of orexigenic signaling systems have been observed in vagal afferents of lean animals. Altered basal cannabinoid (CB₁) receptor expression in the nodose ganglia in obesity has been reported. Whether altered body weight or a high fat diet modifies independent or additive changes in CB₁ expression is unknown. We investigated the expression of CB₁ and orexin 1 receptor (OX-1R) in the nodose ganglia of rats fed ad libitum or food deprived (24 h), maintained on low or high fat diets (HFD), with differing body weights. Male Wistar rats were fed chow or HFD (diet-induced obese; DIO or diet-resistant; DR) or were body weight matched to the DR group but fed chow (wmDR). CB₁ and OX-1R immunoreactivity were investigated and CB₁ mRNA density was determined using in situ hybridization. CB₁ immunoreactivity was measured in fasted rats after sulfated cholecystokinin octapeptide (CCK8s) administration. In chow rats, fasting did not modify the level of CB₁ mRNA. More CB₁ immunoreactive cells were measured in fed DIO, DR and wmDR rats than chow rats; levels increased after fasting in chow and wmDR rats but not in DIO or DR rats. In HFD fasted rats CCK8s did not reduce CB₁ immunoreactivity. OX-1R immunoreactivity was modulated by fasting only in DR rats. These data suggest that body weight contributes to the proportion of neurons expressing CB₁ immunoreactivity in the nodose ganglion, while HFD blunts fasting-induced increases, and CCK-induced suppression of, CB₁-immunoreactivity.

1. Introduction

Energy balance is regulated by the complex interaction of numerous pathways (Bernoud et al., 2011). In the periphery the gastrointestinal tract, liver, pancreas and adipose tissue signal the nutritive status of the body to the central nervous system (CNS) through neuronal pathways and circulating hormones. The CNS and gastrointestinal tract are linked by the bidirectional, hormonal and neuronal pathway termed the gut brain axis. Information from enteroendocrine cells and the enteric nervous system is relayed to the brain stem via vagal afferent neurons, whose cell bodies lie in the nodose ganglia. Plasticity of the nervous and endocrine systems known to be involved in the regulation of energy balance by experiments using CB₁ receptor antagonists which

⁎ Corresponding author at: Department of Physiology & Pharmacology, University of Calgary, 3330 Hospital Drive NW, Calgary, Alberta T2N 4N1, Canada. Tel.: +1 403 220 4601; fax: +1 403 283 2700.
E-mail address: ksharkey@ucalgary.ca (K.A. Sharkey).

© 2013 Elsevier B.V. All rights reserved.
do not pass the blood brain barrier. With their actions restricted to the periphery, URB447 (LoVerme et al., 2009) and AM6545 (Cluny et al., 2010; Randall et al., 2010) have been shown to inhibit food intake in rodents. Furthermore, elevated endocannabinoid levels in the duodenum, liver and pancreas have been found in obese rodents (Izzo et al., 2009). Also, levels of CB1 receptor mRNA are elevated in the stomach of DIO mice (Di Marzo et al., 2008) and nucleus of the solitary tract of obese rats (Bello et al., 2012), and the fasting of lean rats elevates anandamide levels in the duodenum (Gómez et al., 2002).

Recently, it has been reported that vagal afferent CB1 expression in ad libitum fed DIO-leptin resistant and leptin receptor gene-deficient, obese Zucker rats is elevated compared to lean controls and that this is due to the leptin-deficient status of these models (De Lartigue et al., 2012). In other studies, anandamide levels in the small intestine of DIO mice were significantly lower than in their lean counterparts while anandamide levels in the duodenum of obese Zucker rats were increased compared to lean wild-type rats (Izzo et al., 2009). This implies that increased body weight alone may not be responsible for modified endocannabinoid levels in the small intestine and that in obese rodents the ingestion of a high fat diet may have differing actions to those seen with genetic leptin signaling deficiency.

We hypothesized that body weight and a high fat diet would have differing effects on CB1 receptor expression in vagal afferents. The changes in level of expression of cannabinoid CB1 protein and mRNA were determined in the nodose ganglia of rats in differing states of obesity during ad libitum fed and fasted conditions. We also investigated the effects on the protein expression of another orexigenic-mediating receptor, OX-1R.

2. Materials and methods

2.1. Animals

Male Wistar rats, purchased from Charles River (St-Constant, Quebec, Canada), were singly housed, under a 12 h light–dark cycle (lights off 19:00), in sawdust floor, plastic cages and allowed free access to tap water and either standard laboratory chow (SP14-Prolab RMH 2500, 12% kcal from fat, 60% kcal from carbohydrate and 28% kcal from protein; Lab Diet, St. Louis, MO, USA) or a high fat diet (D12451, 45% kcal from fat, 35% kcal from carbohydrate and 20% kcal from protein; Research Diets Inc., New Brunswick, NJ, USA) unless otherwise stated. Rats used in in situ hybridization studies were singly housed in wire-bottom cages suspended above absorbent paper and fed standard laboratory chow. Male Wistar rats (210–230 g, 7–8 weeks old) used in the fasting time-course studies and CCK8s studies were fed standard laboratory chow or a high fat diet (824053, 45% kcal from fat, 35% kcal from carbohydrate and 20% kcal from protein; Special Diet Services, UK). All animal care and experimental procedures were in accordance with the guidelines of the Canadian Council on Animal Care or were conducted under the UK Home Office authority, with approval from the appropriate institutional authorities.

2.2. Experimental groups

Age-matched rats (202.4 ± 1.8 g, 7 weeks old) were randomly assigned to be fed either a standard laboratory chow (n = 6; body weight 201.3 ± 5.6 g) or a high fat diet (n = 20; body weight 202.7 ± 1.8 g; Research Diets Inc. diet). It has been shown that Wistar rats fed a high fat diet (45% calories from fat) for 6 weeks display characteristics of obesity including a greater body weight, insulin resistance, increased fasting plasma insulin, glucose and leptin levels and increased triglycerides in skeletal muscle and plasma (Buettner et al., 2000). The rats in the current study were fed the high fat diet for 11 weeks in order for the DIO and DR phenotypes to develop. The body weight of each rat was measured weekly. After 11 weeks of ingesting a high fat diet the rats were designated as being diet-induced obese (DIO) or diet-resistant (DR) based on their body weight. Rats weighing ≤590 g were classed as DR (n = 9) and those weighing ≥612 g were classed as DIO (n = 11). Other rats were body weight matched (but not age-matched) to the DR group and fed only a chow diet (wmdR; n = 6).

2.3. Tissue collection for immunohistochemistry

Rats were fed ad libitum or fasted for 24 h before being deeply anaesthetized with sodium pentobarbital (>65 mg/kg) administered intraperitoneally (i.p.). Rats were then perfused intracardially with ice-cold phosphate buffered saline (PBS; 1 l/kg rat), to remove the blood, followed by ice-cold 4% paraformaldehyde (PFA; 11 kg rat), to fix the nodose ganglia. The nodose ganglia were microdissected and immersion fixed in 4% PFA for 1 h at room temperature. Tissues were washed in PBS 3 times at 10 min intervals before being cryoprotected overnight in 20% sucrose at 4 °C. The nodose ganglia were then embedded in an optimal cutting temperature medium and exhaustively cut (7 μm) longitudinaly using a cryostat. Sections were sequentially placed on 10 slides. In other experiments, a time-course of fasting was investigated: rats (210–230 g) were fed a high fat diet (Special Diet Services diet) for 4 weeks before being killed by a rising concentration of CO2 after a fasting period of 0, 6, 12, 24, 36 and 48 h (n = 4–5 per time point). The mid and caudal regions of the nodose ganglia were dissected and fixed in 4% PFA.

2.4. CCK8s administration

Rats fed with chow or high fat diet (Special Diet Services diet) for 4 weeks were fasted for up to 24 h. Animals received an i.p. injection of sfulated cholecystokinin octapeptide (CCK8s; 10 nmol dissolved in saline; Bachem, St. Helens, Merseyside, UK) at the end of the fasting period. Rats (n = 3 per group) were killed by a rising concentration of CO2 after a fasting period of 0, 1, 4 and 8 h after receiving CCK8s. The mid and caudal regions of the nodose ganglia were dissected and fixed in 4% PFA.

2.5. Immunohistochemistry

The nodose ganglia sections from chow, DIO, DR and wmdR rats were washed in PBS containing 0.1% Triton X-100 (PBS-T; 3 × 10 min) and then incubated in normal donkey serum (1:10; Jackson ImmunoResearch Laboratories Inc., West Grove, PA) for 30 min at room temperature. Sections were incubated in either rabbit anti-CB1-C-terminal (CB1-CT; 1:1000 (Hajos et al., 2000)), rabbit anti-CB1-N-terminal (CB1-NT; 1:1000) (Tsou et al., 1998; Berghuis et al., 2007) or rabbit anti-OX-1R (1:1000; OX1R11-A; Alpha Diagnostic International, San Antonio, TX) primary antibodies for 48 h at 4 °C. Sections were washed in PBS-T (3 × 10 min) and incubated, at room temperature for 1 h in the secondary antibody donkey anti-rabbit CY3 (1:100; #711-165-152, Jackson ImmunoResearch Laboratories Inc., West Grove, PA, USA). Sections were washed in PBS (3 × 10 min) and mounted in bicarbonate-buffered glycerol before being viewed using a Zeiss Axiosplan fluorescence microscope (Carl Zeiss, Jena, Germany) with images captured using a Retiga 2000R digital camera (QImaging, Surrey, BC, Canada). Four images were captured at 40× magnification from one representative section of the nodose ganglion per slide. The number of immunoreactive cells was determined for each image using Image Pro-Plus software version 7.0 (MediaCybernetics, Silver Spring, MD, USA). For each antibody, the range of intensity was manually determined in preliminary experiments to detect positively stained cells and to exclude background staining. For all analyses, regardless of the immunoreactivity under investigation, the area of intensity detected as a region of immunoreactivity was set to include cell bodies only. The number of immunoreactive cells per nodose ganglion was calculated as a percent of the total cell count. Specificity of the CB1-NT and OX-1R antibodies was tested using immunizing peptides (OX-1R: OX1R11-P, Alpha Diagnostic International). Immunoreactivity to CB1-NT (Tsou et al., 1998) or OX-1R was not observed following incubation with the corresponding peptide. Specificity of the CB1-NT receptor, OX-1R.
and CB1-CT antibodies has previously been demonstrated by a lack of immunoreactivity in brain tissue of CB1 receptor gene-deficient mice (Hu et al., 2012).

Frozen cryostat sections of nodose ganglia from rats involved in the fasting time-course study and the CCK8s study were rinsed in PBS, permeabilized in ethanol, and processed for immunofluorescence as described previously (Burdyga et al., 2004). For detection of CB1, both affinity-purified goat and rabbit polyclonal antibodies were used (1:90; Santa Cruz Biotechnology, Santa Cruz, CA (Burdyga et al., 2004)). The secondary antibodies, FITC-conjugated (Jackson ImmunoResearch Laboratories, West Grove, PA) donkey anti-goat or anti-rabbit IgG, were used as appropriate. Sections were examined using an Axioscan Universal microscope, and images were processed using the Axio Vision 3.0 Imaging system with deconvolution options (Carl Zeiss Vision). The quantification of neurons expressing CB1 was made by counting 400–600 immunoreactive cell profiles in five sections per ganglia, selecting sections separated by 90 μm that passed through the full length of the caudal and mid regions of the ganglia.

2.6. Tissue collection for in situ hybridization

Rats were fed ad libitum (n = 4) or fasted for 24 h (n = 4) before being deeply anesthetized with ketamine (60 mg/kg)/xylazine (7.5 mg/kg, i.p.). They were perfused intracardially with 200 ml of ice-cold isotonic saline followed by 500 ml of 4% PFA. The right nodose ganglia were microdissected and immersion fixed in paraformaldehyde for 7 days. Ganglia were cryoprotected and stored at −30 °C until paraffin inclusion. Fifteen micrometer thick longitudinal sections were cut and placed on microscopic slides.

2.7. In situ hybridization histochemistry

The protocol for in situ hybridization was largely adapted from a previously described technique (Simmons et al., 1989). Deparaffinized sections were successively fixed for 20 min in 4% PFA: this second fixation step improves the hybridization signal, especially when there are low amounts of targeted mRNA. Sections were then digested for 30 min at 37 °C with proteinase K (10 μg/ml in 100 mM Tris–HCl containing 50 mM EDTA, pH 8.0), acetylated with acetic anhydride (0.25% in 0.1 M triethanolamine, pH 8.0), and dehydrated through graded concentrations (50, 70, 95, and 100%) of alcohol. After vacuum drying for at least 2 h, 90 μl of the hybridization mixture, containing an antisense 35S-labeled cRNA probe (107 cpm/ml), was spotted on sections. The sections were successively permeabilized in ethanol, and processed for immunofluorescence as described previously. The DNA templates were treated with 100 U/ml DNase solution containing 0.1 U/ml DNase (Roche Diagnostics), 0.25 mg/ml of tRNA and 50 mM Tris/10 mM MgCl2. The riboprobes were purified using an RNase Mini Kit (Qiagen, Missisauga, ON, Canada), eluted in 150 μl of 10 mM Tris–1 mM EDTA buffer, and incorporated in a hybridization solution containing (per ml) 102 cpm of 35S probe, 52% formamide, 330 mM NaCl, 10 mM Tris (pH 8), 1 mM EDTA (pH 8), 1 × Denhart’s solution, 10% dextran sulfate, 0.5 mg/ml of tRNA, 10 mM DTT, and diethyl pyrocarbonate water. This solution was mixed and heated at 65 °C and then spotted on slides. The specificity of the probe was confirmed by the absence of a positive signal in sections hybridized with sense probe.

2.9. Quantitative analysis of the hybridization signal

The hybridization signals revealed on NTB2-dipped nuclear emulsion slides were examined under dark-field microscopy using an Olympus BX60 microscope (Olympus America, Melville, NY, USA). Images were acquired with a camera (RT Slider, model 2.3.0, Diagnostic Instruments) and analyzed with ImagePro-Plus software version 4.5.1.23. Saturation of the hybridization signal was avoided by adjusting the exposure time for the image with the strongest hybridization signal. The luminosity of the system was set to the maximum, and the saturation warning option was used to visualize saturated regions in the image preview. Thereafter, according to the pixel distribution histogram, the exposure time was adjusted in order to reduce to zero the number of saturated (pure white) pixels. The same exposure time was used throughout the analysis of the entire series. The entire area of the nodose section was outlined; the pixel density of the hybridization signal was measured and then corrected by subtracting background readings taken from areas immediately surrounding the nodose. Four to seven sections were analyzed for each nodose and the average pixel density was considered for statistical analysis.

2.10. Data analysis

Data are expressed as the mean ± s.e.mean and were analyzed using either an unpaired t test or one way analysis of variance followed by Bonferroni’s multiple comparison test where appropriate. p < 0.05 was considered significant.

3. Results

3.1. Body weight

Both groups of rats fed the high fat diet for 11 weeks (DIO: 644.6 ± 8.6 g; DR: 560.9 ± 8.4 g) had a significantly greater (p < 0.01) body weight than the age-matched, chow-fed animals (chow: 498.3 ± 15.5 g; Fig. 1). Furthermore, the body weight of DIO rats was significantly greater (p < 0.001) than DR rats (Fig. 1). DR rats and rats that were body weight matched to DR but fed a chow diet (wmdr): 544.7 ± 9.1 g) had comparable body weights (p > 0.05) and both had a greater body weight (p < 0.05) than the chow-fed control rats (Fig. 1).

3.2. CB1 receptor expression in chow-fed rats

The proportion of cells that were immunoreactive to an antibody raised against the C-terminal of the CB1 receptor was comparable between rats who were ad libitum fed and those who were fasted for 24 h (p > 0.05; Fig. 2A and C). However, the proportion of CB1-NT immunoreactive cells in 24 h fasted rats (CB1-NT cell profile % of the...
total number of cells ± s.e.mean: 21.5 ± 0.7%) was significantly higher than in ad libitum fed chow-fed rats (1.1 ± 0.7%; p < 0.001; Fig. 3). There was no difference in CB1 mRNA density in the nodose ganglia of ad libitum fed vs 24 h fasted rats (p N 0.05; Fig. 2B and D).

3.3. CB1 receptor immunoreactivity

In animals that were ad libitum fed the percentage of CB1-NT immunoreactive cells was significantly higher in DIO (13.3 ± 4.8%; p < 0.05), DR (13.6 ± 4.6%; p < 0.05) and wmDR (8.4 ± 1.8%; p < 0.01) rats compared to chow-fed rats (1.1 ± 0.7%). In DIO and DR rats the percentage of cells that were CB1-NT immunoreactive following a 24 h fast was comparable to the level in the fed ad libitum state (Fig. 3). A time-course of fasting was investigated and, in rats fed a high fat diet for 4 weeks body weight increased from 230 ± 2 g to 432 ± 6g, compared with 233 ± 5 g increasing to 390 ± 9 g in chow-fed controls (p < 0.05). The administration of CCK8s to 24 h fasted, chow-fed rats induced a reduction in CB1 receptor cell profile 1 and 4 h post injection (Fig. 4). Eight hours after the CCK8s injection the percentage of CB1 immunoreactive cells in the nodose ganglia of Chow fasted rats lasted for 24 h had increased back to a comparable level as was observed prior to CCK8s administration (Fig. 4). CCK8s administration had no effect on the CB1 cell profile in the nodose ganglia in rats that were fed a high fat diet and fasted for 24 h (Fig. 4).

3.4. CB1 receptor immunoreactivity following CCK8s administration

In order to examine the effect of a high fat diet on CCK-induced changes in CB1 receptor immunoreactivity rats were maintained on a high fat diet for 4 weeks only which is not long enough for the phenotypes of DIO and DR to develop. In rats fed a high fat diet for 4 weeks body weight increased from 230 ± 2 g to 432 ± 6g, compared with 233 ± 5 g increasing to 390 ± 9 g in chow-fed controls (p < 0.05). The administration of CCK8s to 24 h fasted, chow-fed rats induced a reduction in CB1 receptor cell profile 1 and 4 h post injection (Fig. 4). Eight hours after the CCK8s injection the percentage of CB1 immunoreactive cells in the nodose ganglia of Chow fasted rats lasted for 24 h had increased back to a comparable level as was observed prior to CCK8s administration (Fig. 4). CCK8s administration had no effect on the CB1 cell profile in the nodose ganglia in rats that were fed a high fat diet and fasted for 24 h (Fig. 4).

3.5. OX-1R immunoreactivity

The proportion of OX-1R immunoreactive cells in the nodose ganglia from ad libitum fed DIO (18.4 ± 5.7%), DR (21.3 ± 3.5%) and wmDR (27.8 ± 2.6%) rats was comparable to that in the Chow fed animals (19.8 ± 4.1%; p > 0.05; Fig. 5). There was a significant reduction (p < 0.01) in the percentage of OX-1R immunoreactive cells in fasted DR rats (1.3 ± 1.3%), compared to those fed ad libitum (21.3 ± 3.5%; Fig. 5C). Fasting did not modify the proportion of OX-1R immunoreactive cells in Chow, DIO or wmDR rats (Fig. 5).

4. Discussion

The current study investigated the effects of a high fat diet and body weight on the expression of orexigenic-mediating receptors in the nodose ganglia of rats under ad libitum fed and 24 h fasted conditions. We present data showing that the proportion of N-terminal CB1 receptor immunoreactive cells in the nodose ganglia is dependent on feeding status. Observed differences in immunoreactivity to antibodies raised against differing terminals of the receptor may be indicative of changes...
in receptor activity as opposed to a change in protein expression. The data further suggest that N-terminal CB1 receptor immunoreactivity is enhanced in animals with a greater body weight and that high-fat feeding may blunt fasting-induced increases in receptor expression and CCK-induced reduction in receptor immunoreactivity. This was not observed for OX-1R.

We have confirmed the previous observations (Burdyga et al., 2004; De Lartigue et al., 2012) that immunoreactivity to an antibody raised against the N-terminal of the CB1 receptor is low in the nodose ganglia of ad libitum fed rats maintained on a standard chow diet, and elevated following a period of fasting. However, the present data suggest that the percentage of cells expressing immunoreactivity to an antibody raised against the carboxy tail of the receptor is not reduced when animals are ad libitum fed, and is not modified by fasting. The differences observed in N-terminal immunoreactivity may reflect a change in receptor activity, and that C-terminal immunoreactivity is unchanged may be indicative of unaltered CB1 receptor protein expression in differing feeding statuses.

The cannabinoid CB1 receptor is a class A, rhodopsin-like, G protein-coupled receptor (GPCR) consisting of an extracellular N-terminal, 7 transmembrane helices, with 3 extracellular loops, and an intracellular C-terminal. The C-terminal is involved in G protein binding and thus the induction of signal transduction during receptor activation (Mukhopadhyay and Howlett, 2001; Nie and Lewis, 2001). The N-terminal of the CB1 receptor is not involved in ligand binding or receptor activation but has been suggested to be involved in speculated chaperone-mediated functions such as receptor synthesis and degradation (Andersson et al., 2003). However, upon CB1 receptor activation the N-terminal protein unfolds (Gupta et al., 2007). This suggests that activation-induced unfolding of the N-terminal results in the exposure of the epitope to which the CB1-NT antibody used in the current studies was raised. We show that the proportion of cells displaying immunoreactivity to the N-terminal of CB1, is very low when lean, chow-fed animals are ad libitum fed and that this is increased following a 24 h fast. Such fluctuations were not seen with the degree of immunoreactivity to the C-terminal of the receptor. The specificity of the antibody used in the current studies has been confirmed by a lack of staining in CB1 receptor knockout mice suggesting that the possibility of non-specific staining is highly unlikely. Moreover, our findings suggest that CB1 mRNA expression in the nodose ganglia of chow fed rats is similarly

Fig. 3. CB1-NT immunoreactivity in nodose ganglia from chow fed, weight matched to diet resistant (wmDR), diet resistant (DR) and diet-induced obese (DIO) rats ad libitum fed (ad lib fed; A) or fasted for 24 h (fasted; B). Scale bar: 100 μm. An example of an immunopositive cell (closed arrow) and an immunonegative cell (open arrow) is indicated. Quantitative analysis of CB1-NT immunoreactivity (C) in nodose ganglia from chow fed, weight matched to diet resistant (wmDR), diet-induced obese (DIO), and diet resistant (DR) rats ad libitum fed or fasted for 24 h. Bars represent the mean ± s.e.mean, n = 3–5 per group.* (p < 0.05) and *** (p < 0.001) denote a significant difference to the corresponding ad lib fed group analyzed using unpaired t-test.
not modified by feeding status. CB₁ mRNA transcript was evident at comparable levels in cell bodies of vagal afferents in rats which were fed ad libitum and in those which were fasted for 24 h. While these data are similar to those reported by Jelsing and colleagues (Jelsing et al., 2009) it has also been reported that CB₁ mRNA was reduced by feeding and increased by fasting in the nodose ganglia of rats (Burdyga et al., 2004). Technical differences between the studies may account for the differing results. RNA probes were used in the present study, and also in that reported by Jelsing and colleagues (Jelsing et al., 2009), whereas a three-oligonucleotide probe was used in the study where feeding status-modifications in mRNA density were reported (Burdyga et al., 2004). RNA probes offer a high degree of sensitivity and stability (due to high specific activities, genetic complexities and hybridization energy between RNA–RNA hybrids). Furthermore, higher stringency of the RNA–RNA hybridization results in the detection of less abundant transcripts. The current findings suggest that neither mRNA nor overall protein expression is altered by feeding status, but rather our results using the N-terminal antibody may reflect activated and inactivated states of the receptor. Levels of the endocannabinoid 2-arachidonoylglycerol (2-AG) in rat hypothalamus (Kirkham et al., 2002) and in whole mouse brain (Hanus et al., 2003) have been

Fig. 4. The effect of CCK8s (10 nmol, i.p.) administration on CB₁-NT immunoreactivity in nodose ganglia from 24 h fasted rats fed chow or high fat diet (HFD). Data points represent the mean ± s.e.mean, n = 3 per group.

Fig. 5. OX-1R immunoreactivity in nodose ganglia from chow fed, weight matched to diet resistant (wmDR), diet resistant (DR) and diet-induced obese (DIO) rats ad libitum fed (ad lib fed; A) or fasted for 24 h (fasted; B). Scale bar: 100 μm. An example of an immunopositive cell (closed arrow) and an immunonegative cell (open arrow) is indicated. Quantitative analysis of OX-1R immunoreactivity (C) in nodose ganglia from chow fed, weight matched to diet resistant (wmDR), diet-induced obese (DIO), and diet resistant (DR) rats ad libitum fed or fasted for 24 h. Bars represent the mean ± s.e.mean, n = 3 per group. ** (p < 0.01) denotes a significant difference to the corresponding ad lib fed group analyzed using unpaired t-test.
shown to be significantly increased after a 24 h period of fasting. Furthermore, fasting for 24 h induced an increase in the levels of the major endocannabinoids, 2-AG and anandamide, in rat limbic forebrain (Kirkham et al., 2002), duodenum, pancreas and liver (Izzo et al., 2009) compared to non-food deprived animals. That there are low levels of endocannabinoids during fed states implies that there is little opportunity for CB1 receptor activation. This suggests that the conformation of the N-terminal of the receptor would be such that the antibody directed towards this region could not bind, resulting in low immunoreactivity in the ad libitum fed rat nodose ganglia in the current study. It is suggested that CB1 activation plays a role in the maintenance of fasting-induced neurochemical phenotypes in the nodose ganglia such as increased CB1 and reduced Y2R expression (Burdyga et al., 2010). Administration of a CB1 receptor antagonist/inverse agonist blocked fasting-induced changes in expression of CB1 and the anorexigenic-mediating receptor Y2 in the nodose ganglia (Burdyga et al., 2010). This suggests that CB1 receptors in the nodose ganglia, as opposed to the central or peripheral terminals of the vagal afferents are involved in this effect. In the current study, the percentage of cells that showed CB1 N-terminal immunoreactivity was increased in the nodose ganglia of fasted rats and this may reflect receptor activation by elevated endocannabinoid levels, leading to the maintenance of fasting-induced neurochemical phenotypes in this structure.

We confirmed the development of two rat phenotypes in response to a high fat diet; those with a greater body weight, termed diet-induced obese (DIO) and those which were leaner termed diet-resistant (DR) (Levin et al., 1983). We used these models to examine whether feeding obese (DIO) and those which were leaner termed diet-resistant (DR) rats demonstrated higher levels of endocannabinoids in the duodenum, liver and pancreas compared to lean rats and in the duodenum these levels were increased even further upon fasting (Izzo et al., 2009). This may suggest that the elevated level of endocannabinoid production in response to the two stimuli of high body weight and fasting is additive. As described above, high fat feeding is also a reported stimulus of endocannabinoid production and this may also act additively with fasting and a high body weight to induce endocannabinoid production. This may lead to downregulation of CB1 receptor and thus in the DIO and DR rats no further increase in activity was possible in response to fasting. It may be argued that fasting-induced increases in CB1–NT immunoreactivity in these rats are delayed however, the current studies suggest that even when the fasting period is increased to 48 h immunoreactivity remains comparable to that in the ad libitum fed state.

There is growing evidence that in vagal afferent neurons the actions of CCK can influence the expression of receptors that are involved in the control of food intake, including CB1 (Dockray and Burdyga, 2011). CB1 and CCK are colocalized in neurons in the rat nodose ganglia and CCK8 administration reduces CB1 expression in fasted chow-fed rats (Burdyga et al., 2004). However, the present study suggests that exogenous CCK has no effect on food deprived CB1 receptor immunoreactivity in heavy rats fed a high fat diet, suggesting that control of CCK over this orexigenic signaling pathway may be lost following ingestion of high levels of fat and/or due to a heavier body weight. The level of CCK-1R mRNA in the nodose ganglia is reported to be the same in weight-matched rats fed either chow or a high fat diet for 14 days (Broberger et al., 2001). However, mice fed a high fat diet for 15 days that had a greater body weight than chow fed mice had a lower level of CCK-1R mRNA in the nodose ganglia and in these mice, exogenous CCK was less effective to reduce food intake than in chow fed mice (Nefti et al., 2009). This may suggest that a greater body weight, rather than a high fat diet plays a part in these differences. However, exogenous CCK is more efficacious to reduce food intake in chow fed rats that are obesity prone compared to chow fed rats that are significantly leaner (Swartz et al., 2010) and CCK1R mRNA is elevated in the nodose ganglia of DIO rats yet is unchanged in rats who are resistant to the obesogenic effects of a high fat diet (Paulino et al., 2009), suggesting that body weight and fat content of the diet may both exert modifications in CCK signaling. The studies discussed above also highlight possible differences in response to CCK depending on the animal model used, with the hypophagic effect of exogenous CCK being less efficacious in mice (Nefti et al., 2009) and more efficacious in DIO rats (Swartz et al., 2010). It has been suggested that leptin signaling is essential for endogenous CCK-induced modifications of neurochemical phenotype in cell bodies of the nodose ganglia (De Lartigue et al., 2012): In rats that were fasted for 24 h before being fed for 1 h (a model of meal-induced endogenous CCK release),

For example, in mice fed a high fat diet anandamide, but not 2-AG, levels were reduced in the stomach (Di Marzo et al., 2008) yet elevated in the liver (Osei-Hyiaman et al., 2005). Anandamide and 2-AG levels were increased in the pancreas of DIO mice compared to lean controls, and in visceral fat only 2-AG was increased in DIO mice (Matias et al., 2006). Increased 2-AG but reduced anandamide was observed in the duodenum of mice fed a high fat diet (Izzo et al., 2009). A general increase in endocannabinoid levels was observed in the skeletal muscle, brown adipose tissue, adrenal gland, heart and kidneys in mice fed high fat diets, with the time of onset and duration of the changes being dependent on the fatty acid content of the diet (Matias et al., 2008). CB1 protein expression in purified liver plasma membrane is increased in mice fed a high fat diet (Osei-Hyiaman et al., 2005). A high fat diet did not modify CB1 immunoreactivity in mouse pancreatic islets, visceral or subcutaneous adipose tissue (Starowicz et al., 2008). After being fed a high fat diet CB1 mRNA levels were reduced in mouse stomach antrum (Di Marzo et al., 2008) and in rat cingulate pre-frontal cortex, ventrolateral parcellular paraventricular hypothalamic nucleus and ventromedial hypothalamic nucleus (Timofeeva et al., 2009). Furthermore, obese Zucker fa/fa rats demonstrated higher levels of endocannabinoids in the duodenum, liver and pancreas compared to lean rats and in the duodenum these levels were increased even further upon fasting (Izzo et al., 2009). This may suggest that the elevated level of endocannabinoid production in response to the two stimuli of high body weight and fasting is additive. As described above, high fat feeding is also a reported stimulus of endocannabinoid production and this may also act additively with fasting and a high body weight to induce endocannabinoid production. This may lead to downregulation of CB1 receptor and thus in the DIO and DR rats no further increase in activity was possible in response to fasting. It may be argued that fasting-induced increases in CB1–NT immunoreactivity in these rats are delayed however, the current studies suggest that even when the fasting period is increased to 48 h immunoreactivity remains comparable to that in the ad libitum fed state.

There is growing evidence that in vagal afferent neurons the actions of CCK can influence the expression of receptors that are involved in the control of food intake, including CB1 (Dockray and Burdyga, 2011). CB1 and CCK are colocalized in neurons in the rat nodose ganglia and CCK8 administration reduces CB1 expression in fasted chow-fed rats (Burdyga et al., 2004). However, the present study suggests that exogenous CCK has no effect on food deprived CB1 receptor immunoreactivity in heavy rats fed a high fat diet, suggesting that control of CCK over this orexigenic signaling pathway may be lost following ingestion of high levels of fat and/or due to a heavier body weight. The level of CCK-1R mRNA in the nodose ganglia is reported to be the same in weight-matched rats fed either chow or a high fat diet for 14 days (Broberger et al., 2001). However, mice fed a high fat diet for 15 days that had a greater body weight than chow fed mice had a lower level of CCK-1R mRNA in the nodose ganglia and in these mice, exogenous CCK was less effective to reduce food intake than in chow fed mice (Nefti et al., 2009). This may suggest that a greater body weight, rather than a high fat diet plays a part in these differences. However, exogenous CCK is more efficacious to reduce food intake in chow fed rats that are obesity prone compared to chow fed rats that are significantly leaner (Swartz et al., 2010) and CCK1R mRNA is elevated in the nodose ganglia of DIO rats yet is unchanged in rats who are resistant to the obesogenic effects of a high fat diet (Paulino et al., 2009), suggesting that body weight and fat content of the diet may both exert modifications in CCK signaling. The studies discussed above also highlight possible differences in response to CCK depending on the animal model used, with the hypophagic effect of exogenous CCK being less efficacious in mice (Nefti et al., 2009) and more efficacious in DIO rats (Swartz et al., 2010). It has been suggested that leptin signaling is essential for endogenous CCK-induced modifications of neurochemical phenotype in cell bodies of the nodose ganglia (De Lartigue et al., 2012): In rats that were fasted for 24 h before being fed for 1 h (a model of meal-induced endogenous CCK release),
fasting-induced elevation of CB1 receptor expression in the nodose ganglia of DIO animals was not down-regulated by feeding and it was suggested that this was due to the leptin resistant status of the rats (De Lartigue et al., 2012). It has been reported that after 8 weeks on a high fat diet vagal afferent neurons of DIO but not DR rats exhibited leptin resistance (De Lartigue et al., 2011). In the present study, animals were fed a high fat diet for only 4 weeks and the DR vs DIO phenotype had not presented; thus it is unlikely that those rats were leptin resistant compared to the chow-fed controls. Our findings suggest that changes in CCK signaling in rats fed a high fat diet, which were heavier than chow-fed controls, may result in a loss of CCK-induced inhibition of CB1 receptor immunoreactivity, and thus maintain the higher levels of basal CB1 activity we observed in fed ad libitum rats. The data suggest that the ingestion of a high fat diet or a heavier body weight alters the ability of CCK to negatively influence endocannabinoid signaling and this may play a role in the maintenance of a positive energy balance.

The CB1 immunoreactivity changes in the nodose ganglia observed in response to feeding status, high fat feeding and body weight were not similarly seen in OX-1R immunoreactivity in the current study. In the ad libitum fed state immunoreactivity to OX-1R was present and a 24 h fast did not modify this in the nodose ganglia of Chow fed rats. OX-1R gene expression in lean rat nodose ganglia was not altered by fasting (Burdyga et al., 2004) and the present study suggests that protein expression is similarly not altered by food deprivation in this structure. Furthermore we present data suggesting that the proportion of OX-1R-immunoreactive cells in DIO rats and heavier Chow-fed rats was comparable to leaner Chow-fed animals and this was not modified by feeding status. A high fat diet had no effect on hypothalamic OX-1R mRNA levels (White et al., 2005) and the current study suggests that high fat feeding does not alter OX-1R protein in the nodose ganglia. However, it would appear that in animals resistant to the obesogenic effects of high fat feeding, food deprivation induces a down-regulation in the number of OX-1R receptor immunoreactive cells. This was not observed in weight-matched rats that were fed a standard Chow diet and thus suggests that this occurs in diet resistant animals only when exposed to a high fat diet. This reduction in orexin-1 receptor expression may play a role in the generation of the leaner phenotype observed in these rats.

5. Conclusion

These results suggest that a heavier body weight may contribute to an elevated proportion of neurons expressing CB2-NT immunoreactivity in the nodose ganglion of rats fed ad libitum and that a high-fat diet blunts the increase of CB2-NT immunoreactivity in response to fasting, and also suppresses the inhibitory action of CCK on CB1 immunoreactivity in the nodose ganglia. This indicates that the two parameters of a greater body weight and high fat feeding may act in differing ways but with the same outcome: that CB1 activation in the nodose ganglia is elevated which may contribute to a physiologically-inappropriate stimulation of food intake, leading to the maintenance of obesity.

Disclosures

The authors disclose no conflicts of interest.

Acknowledgments

The authors thank Winnie Ho for the technical assistance. These studies were supported by the grants from the Canadian Institutes of Health Research (Team grant in the Neurobiology of Obesity: no. OTG-183738) (K.A.S. and D.R.), the Biotechnology and Biological Sciences Research Council (G.C.J.D. and G.B.) and the NIH (DA01322 and DA021696) (K.M.). K.A.S. is an AHFMR Medical Scientist and the Crohn’s and Colitis Foundation of Canada Chair in Inflammatory Bowel Disease Research. N.L.C. is an Alberta Innovates-Health Solutions (AI-HS) fellow.

References


