Monounsaturated Fatty Acids Prevent the Aversive Effects of Obesity on Locomotion, Brain Activity, and Sleep Behavior

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Fat and physical inactivity are the most evident factors in the pathogenesis of obesity, and fat quality seems to play a crucial role for measures of glucose homeostasis. However, the impact of dietary fat quality on brain function, behavior, and sleep is basically unknown. In this study, mice were fed a diet supplemented with either monounsaturated fatty acids (MUFAs) or saturated fatty acids (SFAs) and their impact on glucose homeostasis, locomotion, brain activity, and sleep behavior was evaluated. MUFAs and SFAs led to a significant increase in fat mass but only feeding of SFAs was accompanied by glucose intolerance in mice. Radiotelemetry revealed a significant decrease in cortical activity in SFA-mice whereas MUFAs even improved activity. SFAs decreased wakefulness and increased non–rapid eye movement sleep. An intracerebroventricular application of insulin promoted locomotor activity in MUSA-fed mice, whereas SFA-mice were resistant. In humans, SFA-enriched diet led to a decrease in hippocampal and cortical activity determined by functional magnetic resonance imaging techniques. Together, dietary intake of MUFAs promoted insulin action in the brain with its beneficial effects for cortical activity, locomotion, and sleep, whereas a comparable intake of SFAs acted as a negative modulator of brain activity in mice and humans.

Besides a decrease in physical activity, the main factor in the progression toward insulin resistance, obesity, and type 2 diabetes is overeating and, in particular, dietary fat intake. Although the daily intake of dietary fat remained stable over the past decades, the number of patients with obesity and type 2 diabetes reached epidemic proportions worldwide. A large number of studies blamed both the quantity and quality of dietary fat as a major culprit impairing insulin sensitivity and insulin secretion, but recent studies suggested that fat quality more strongly correlated with weight gain and metabolic alterations than fat quantity (1–3). In particular, dietary monounsaturated free fatty acids (MUFAs) prevent or ameliorate metabolic syndrome and type 2 diabetes, and a fatty acid pattern associated with insulin resistance is characterized by high levels of SFAs and a low concentration of the unsaturated n-6 fatty acid linoleic acid (4, 5). By magnetoencephalography studies in humans, we recently reported that a high-fat diet based on lard mediates insulin resistance in the mouse brain and impairs cortical activity and locomotion, which might further promote glucose intolerance, physical inactivity, and obesity (16), however, the diet used in that study contained a large amount of fat. Although the aversive effects of SFAs overload on glucose metabolism are well documented in a number of studies (17), the impacts of MUFAs on insulin sensitivity in the brain and brain activity are largely unknown. The current design therefore reflects variations of fat quality as present in Western societies and consisted of two moderate, isocaloric, fat-enriched diets that just differ in fat quality but not quantity. Canola oil is rich in MUFAs, with >63% oleic acid, and low in SFAs. By contrast, the fat
content of milk fat is dominated by SFAs, with a high content of palmitic acid (for analysis of fat quality, please see Supplementary Fig. 1 and Supplementary Table 1).

In contrast to an isocaloric, SFA-enriched diet, we report that the intake of an MUFA-enriched diet protected from obesity-associated glucose intolerance and brain and physical inactivity. Further, an intervention study in humans demonstrated that SFA intake deteriorates brain activity in the hippocampus and the inferior parietal cortex whereas MUFAs did not display aversive effects.

RESEARCH DESIGN AND METHODS

Animals. Male C57BL/6J mice were purchased from Charles River Laboratories (Sulzfeld, Germany). Four-week-old animals were fed Chow diet (5% total fat by weight) or Chow diet supplemented with either 3.3% fat by weight from canola oil or milk fat (Altromin, Lage, Germany) for 8 weeks, and diet feeding continued during electrocorticography (ECoG) measurements. A detailed analysis of the fatty acid composition of the diets is given in Supplementary Fig. 1 and Supplementary Table 1. All animal procedures were approved by local government authorities for animal research according to the guidelines of laboratory animal care.

Glucose and insulin tolerance tests. Overnight-fasted mice were intraperitoneally injected with 2 g/kg body weight of α-g-glucose, and blood samples from tail bleeds were drawn for the determination of glucose using a Glucometer Elite (Bayer, Elkhart, IN) at basal and 15, 30, 60, and 120 min after injection. Insulin tolerance testing was performed with 1 unit/kg body weight of human insulin intraperitoneally in fed mice (Novo Nordisk, Bagsvaerd, Denmark) at 8:00 a.m., and serum insulin was measured at basal and blood glucose levels at basal and 15, 30, and 60 min after injection.

ECoG measurements and locomotor activity. Each mouse received an implantable telemetry electrocorticography transmitter and was instrumented with a guide cannula for microinjection of substances into the lateral ventricles as described previously (16). Telemetry signals were recorded and stored digitally using the Dataquest A.R.T. 4.2 software (DSI). Data analysis for ECoG measurements was performed as described previously (18). Basal ECoG measurements were conducted for four consecutive days.

Intracerebroventricular injection of human insulin or palmitic acid. Human insulin was delivered in a volume of 2 μL over 1 min into fasted mice intracerebroventricularly. The mice received an injection of either human insulin (3.75 μIU total, purchased from Novo Nordisk) or vehicle (0.9% NaCl) in random order 4 days apart. Telemetry signals were recorded continuously for 240 min postinjection. Chow-fed control mice were injected intracerebroventricularly with vehicle or palmitic acid (225 μM) total shortly before the darkness period started (7:00 P.M.).

Sleep recordings and analysis of sleep data. ECoG signals were acquired digitally by using Somnologica Science software (version 3.3.1; Medical, Reykjavík, Iceland). The three vigilance states, wake, rapid-eye-movement (REM) sleep, or non-REM (NREM or slow-wave) sleep, were scored in 4-s epochs.

Clinical chemistry. Serum levels of insulin, adiponectin, and leptin were obtained by RIA (Millipore, Billerica, MA). Analysis of nonesterified fatty acid (NEFA), cholesterol, triglycerides, and lipoproteins was performed on the automated clinical chemistry analyzer (ADVIA 1800; Siemens Healthcare Diagnostics, Eschborn, Germany).

Western blot analysis. Overnight-fasted mice received a bolus of human insulin (1 unit/mouse i.v. for 10 min; Novo Nordisk). Brain and liver tissues were homogenized as previously described (19). Lysates containing equal amounts of protein were immunoprecipitated with antibodies against insulin receptor (IR) (20) or IR substrate 2 (IRS2) (Millipore, Schwalbach, Germany). Antibodies directed against phosphotyrosine PY-20 (Santa Cruz, La Jolla, CA) or IRS2 (Millipore, Schwalbach, Germany) were used for Western blot analysis, one-way ANOVA was used to analyze the main effect of fat-enriched diet by using the difference between the normalized whole-brain fractional amplitude of LFF maps of the pre- and postmeasurement. A P value of <0.001 was considered statistically significant. To compare the effect of SFA- and MUFA-enriched diet with the control diet, one-way ANOVA with Tukey post hoc test for multiple comparison analysis was used. Significance was determined at P < 0.05.

Statistical analysis. Data are expressed as mean ± SEM of the indicated number of experiments. All data were analyzed using Origin 8.1, and significance was performed using Student t test and one-way ANOVA with Dunnett or Bonferroni post hoc test for multiple comparison analysis. Significance was set at P < 0.05. Western blot results are presented as direct comparison of bands in autoradiographs and quantified by optical densitometry (Herolab, Wiesloch, Germany).

RESULTS

SFA-enriched diet leads to more weight gain upon isocaloric food intake compared with MUFAs. To evaluate the metabolic consequences of fat quality, 4-week-old mice were weaned on MUFA- or SFA-enriched diets and compared with Chow-fed animals. As early as 4 weeks on a fat-enriched diet, both groups displayed a significant increase in body weight (Fig. 1A), and the SFA-fed mice were significantly heavier than the MUFA-fed group (Fig. 1B).

However, there was no difference in the total food intake in grams (data not shown), as well as in the daily calorie intake between the fat groups (Fig. 1C).

SFA-enriched diet promotes glucose intolerance by inhibiting insulin secretion and impairs insulin sensitivity. Intraperitoneal glucose tolerance tests (GTTs) revealed that SFA-fed mice were glucose intolerant (Fig. 1D) compared with the MUFA-enriched diet and Chow group, and fasted blood glucose tended to be higher in the SFA group; however, it did not reach significance (P = 0.06). Serum insulin concentrations during the GTT showed that glucose-stimulated insulin secretion was impaired in the
FIG. 1. Metabolic consequences of fat-enriched diets in C57BL/6 mice. C57BL/6 mice were weaned on MUFA-enriched (gray), SFA-enriched (black), or chow (white) diet for 8 weeks. A and B: SFA- and MUFA-fed mice displayed significantly increased body weight gain after 6 weeks compared with the control group (n = 15–30/group; *P < 0.05, ***P < 0.001). C: Food intake in kcal/day for control (white circles), MUFA (gray circles), and SFA (black circles) groups during the feeding period. Asterisks mark significance to controls (***P < 0.001 and **P < 0.005).

D: Intraperitoneally injected glucose during a GTT was cleared slower and less effectively in SFA-fed mice (black circles) compared with controls (white circles) and MUFA-fed animals (gray circles) (n = 15–30/group; *P < 0.05 and **P < 0.005 to controls; #P < 0.05 to MUFA group). E: Attenuated serum insulin concentration during the GTT in SFA-fed mice (black circles). Increased fed (F) and fasted (G) serum insulin concentrations in the SFA-diet animal group (*P < 0.05 and **P < 0.005). H: Blood glucose concentrations from tail bleed during an intraperitoneal insulin tolerance test in control (white), MUFA (gray), and SFA (black) mice (n = 15–30/group; *P < 0.05 SFA group to controls). Data are presented as mean ± SEM.
SFA group whereas MUFA enrichment did not affect insulin secretion (Fig. 1E). The fed insulin concentrations were elevated in both fat groups (Fig. 1F) whereas fasted insulin was solely increased in the SFA group (Fig. 1G). Glucose concentrations in the fed state were significantly increased in the SFA group, and feeding with both diets did not reveal a significant change in whole-body insulin sensitivity, determined by an intraperitoneal insulin tolerance test (Fig. 1H).

NEFA, triglyceride, and leptin concentrations were increased by SFA-enriched diet, whereas lipoproteins, cholesterol, and adiponectin levels were not affected. A significant difference was observed for NEFA levels between SFA-fed and control mice (Fig. 2A), and triglyceride concentrations were significantly increased both in MUFA- and SFA-fed animals (Fig. 2B). Both fat-enriched diets did not alter total cholesterol and lipoproteins, such as HDL and LDL (Fig. 2B). Although there was a significant increase in serum leptin concentrations by SFA enrichment (Fig. 2C), adiponectin levels were not affected by either diet (Fig. 2D).

**MUFA- and SFA-fed mice are characterized by increased total and visceral adipose tissue.** Both fat-enriched diets led to a significant increase in visceral and total adipose tissue as determined by MRI measurements compared with the chow-fed animals, whereas the SFA-fed mice displayed the highest percentage of visceral and total fat mass (Fig. 2E).

Insulin-mediated phosphorylation of the IR and AKT is attenuated in liver tissue of SFA-fed mice. As increased fasting insulin levels in mice fed an SFA diet suggested hepatic insulin resistance, we determined insulin action in liver tissues after insulin injection. Thereby,
insulin significantly increased tyrosine phosphorylation of IR in chow-fed mice (Fig. 3A) and to a lesser degree in MUFA-fed animals, whereas SFA-fed mice displayed insulin resistance. At the level of AKT, insulin promoted phosphorylation of AKT in the chow- and MUFA-fed group, whereas insulin was not able to increase phosphorylation of AKT in SFA-fed mice (Fig. 3B).

**SFA enrichment impairs insulin sensitivity in the brain and lowers cerebrocortical activity.** Insulin was able to increase phosphorylation of IR in brain tissues in the chow group and to a lesser degree in the MUFA-fed group whereas the SFA-fed mice displayed insulin resistance (Fig. 4A). However, tyrosine phosphorylation of IR and IRS2 was enhanced in the basal state (Fig. 4A) in both fat-enriched diet groups compared with the control group. Insulin resistance in the brain of SFA-fed mice was further supported by the fact that insulin-mediated phosphorylation of AKT (S473) (Fig. 4A) was greatly diminished in this group. Interestingly, hyperinsulinemia in the SFA-fed mice led to an enhanced basal phosphorylation of AKT, which was absent in the MUFA- and chow-fed group. In addition, cortical activity was measured. As depicted in Fig. 4B, it became evident that mice fed an SFA-enriched diet were characterized by a decline in cortical activity in all of the frequency bands. Of note, MUFA-fed mice showed an even elevated cortical activity and suggested that MUFAs might even have a beneficial effect on the level of cortical activity.

Previously, we demonstrated that insulin given intracerebroventricularly promotes locomotor activity in lean mice whereas diet-induced obese mice were resistant. To translate insulin-mediated modulation of cortical activity to behavior, locomotion in freely moving mice after intracerebroventricular injection of insulin was monitored. An acute insulin application resulted in less pronounced locomotion in SFA-fed mice compared with MUFA- or chow-fed animals (Fig. 4C), suggesting that SFA enrichment negatively affects insulin sensitivity in the brain on the behavioral level.

**Different sleep-wake patterns in MUFA- and SFA-fed mice and after intracerebroventricular application of palmitic acid.** Based on recent data on metabolism and sleep and in the face of altered brain activity in mice fed a MUFA- or SFA-enriched diet, we tested whether the fat-enriched diets might lead to an altered or even disturbed sleep-wake pattern in mice. All mice exhibited a clear-cut diurnal sleep-wakefulness rhythm, and all mice spent more time awake in the 12-h darkness period than in the 12-h daylight period (Fig. 5A), but SFA-fed mice were significantly less awake during the dark period. Interestingly, SFA-enriched diet had a significantly stronger effect on NREM sleep than on REM sleep in the dark period (Fig. 5B and C). Moreover, fat diet led to a significant increase in NREM sleep in the presence of decreased wakefulness compared with the control group (Fig. 5B). MUFA-fed mice spent 19% less time in REM sleep during the daylight period compared with controls and 24% less compared with SFA-fed mice (Fig. 5C). To prove the fact that SFAs indeed alter wakefulness and sleep behavior, we injected palmitic acid intracerebroventricularly into chow-fed mice shortly before the dark period. Thereby, palmitate significantly decreased the time awake during the night phase while it increased NREM sleep, as demonstrated for the SFA-fed mice (Fig. 5D and E). As expected, palmitic acid had a minor effect on REM sleep (Fig. 5F), as was demonstrated in the SFA-fed mice (Fig. 5C).

**Body weight–matched SFA-fed mice are glucose intolerant and characterized by increased fasted serum insulin levels and greatly impaired cortical activity compared with MUFA-fed animals.** To finally evaluate the impact of certain dietary fat on metabolic parameters and brain activity independent of body weight, body weight–matched mice (MUFA 27.2 ± 0.6 g, SFA 27.4 ± 0.8 g; control 27.0 ± 0.6 g; n = 4 per group) were compared. Those that received the SFA-enriched diet were glucose intolerant (Fig. 6A) compared with the MUFA-enriched diet and mice of the control group, and this was accompanied by a slightly increased fasted blood glucose concentration (P = 0.05) (Fig. 6B). Further, serum insulin concentrations during the GTT were even enhanced by MUFA enrichment (Fig. 6C), whereas body weight–matched, SFA-fed animals tend to have increased fasted insulin concentrations (P = 0.13) (Fig. 6D). This was further supported by an enhanced homeostasis model assessment of insulin resistance in the SFA-fed group, demonstrating insulin resistance (Fig. 6E). Strikingly, basal ECoG analysis revealed a significant decrease in overall cortical activity in all of the frequency bands in the SFA-fed group, indicating that the higher content of SFAs has a negative impact on neuronal activity in the brain, whereas MUFA enrichment induced only a slight decline in the alpha and beta frequency band (Fig. 6F).

**Body weight–matched SFA-fed animals displayed insulin resistance in the brain.** To validate the impact of distinct dietary fatty acids on neuronal insulin sensitivity in vivo, ECoG was analyzed in the postinjection period after an acute intracerebroventricular insulin application. Clearly, intracerebroventricular insulin injection was accompanied by an increase in the delta, theta, alpha, and beta frequency bands as previously described (16), and this was also true for MUFA-fed animals (Fig. 7A–D), whereas SFA enrichment led to a significant decrease in cortical activity compared with body weight–matched controls, as

![Figure 3](image-url)
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FIG. 4. Influence of SFA and MUFA enrichment on insulin sensitivity in the brain and the impact on cortical and locomotor activity. A: Attenuated tyrosine phosphorylation (PY) of IR, IRS2, and phosphorylation of AKT (S473) as well as protein expression of AKT was analyzed by SDS-PAGE after intravenous insulin injection in overnight-fastened MUFA-, SFA-, or chow-fed animals. Densitometric quantification of PY-IR (relative increase of PY-IR compared with insulin-stimulated control mice; n = 5–6/group; *P < 0.05) after intravenous insulin (cross-hatched bars) or vehicle (white bars) injection is presented in histograms. A representative blot is shown. B: Analysis of basal power spectral density (indicated for delta [0.5–4 Hz], theta [4–8 Hz], alpha [8–12 Hz], and beta [12–30 Hz] frequencies) using fast Fourier transformation and estimated by ECoG in MUFA- (gray bars), SFA- (black bars), and chow-fed (white bars) animals. Basal ECoG measurements were conducted during a 4-day period (n = 11–24/group; *P < 0.05, **P < 0.005, and ***P < 0.001 compared with controls; ###P < 0.005 and ####P < 0.001 between fat-enriched diet groups). C: An acute intracerebroventricular insulin application (cross-hatched bars) vs. vehicle (white bars) results in less pronounced locomotor activity in SFA-fed mice. ***P < 0.001 and *P < 0.05 to vehicle. Data are mean ± SEM. Ins, insulin.

**DISCUSSION**

The scope of this study was to characterize the effect of a moderate, isocaloric fat diet either supplemented by MUFAs (canola oil) or by SFAs (milk fat), which only differed in fat quality but not quantity. Measures of glucose homeostasis, lipid metabolism, brain activity, and sleep architecture were performed to judge the value of fat quality in the mouse brain, and this was verified in humans.

We previously showed that feeding of mice with high-fat diet resulted in impaired insulin action in the brain (16). Because of the extreme fat load rich in SFAs, the current study focused on the impact of a moderate, isocaloric fat enrichment with diets that differ in their MUFA and SFA content. Of note, both fat-enriched diets contained less polyunsaturated fatty acids compared with chow and further studies are needed to address this issue in more detail.

We show that slightly elevated, chronic SFA intake impaired peripheral glucose metabolism and led to insulin resistance, which was also verified in body weight–matched animals. Although MUFA-fed animals gained more body weight and fat mass compared with controls, they did not show signs of glucose intolerance. This is in agreement with a former study in humans where an increase in MUFAs augmented glucose-stimulated insulin secretion whereas SFAs were less effective (24). Moreover, the MUFA-enriched diet protected mice from insulin resistance, both detected in their MUFA and SFA content. Of note, both fat-enriched diets contained less polyunsaturated fatty acids compared with chow and further studies are needed to address this issue in more detail.

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well as MUFA-fed animals (Fig. 7A–D). Taken together, these data further demonstrate that insulin failed to stimulate neuronal activity in body weight–matched SFA-fed mice independent of body weight.

**fMRI before and after SFA and MUFA enrichment in healthy human subjects.** Our rodent data provide evidence for SFA-mediated insulin resistance in the brain; however, the relevance in humans is hard to predict. Recently, we demonstrated that brain function during rest is altered in obese subjects (13). Although fasting insulin levels were associated with brain activity in the cortex and the putamen, the impact of dietary fat on networks that represent object recognition and food regulation are unknown. To proceed in this respect, we made use of a well-established approach where we measured brain activity by fMRI after a moderate SFA or MUFA enrichment in healthy human subjects. Before the intervention, there was no significant difference of fasting insulin (pmol/L) between the groups (one-way ANOVA, P > 0.05) (SFA enriched 38.6 ± 16.2, MUFA enriched 41.4 ± 17.7, or control group 51.1 ± 23.2). Upon fat intake, there was no significant increase in body weight, blood glucose, or lipid levels (data not shown). Dietary protocols revealed no significant difference in caloric intake (kcal/day) during the intervention between the three groups (one-way ANOVA, P > 0.05) (SFA enriched 2425 ± 665, MUFA enriched 2287 ± 571, or control group 2060 ± 309). All subjects underwent a resting-state fMRI before and after the 12 weeks of fat consumption. There was a significant main effect in whole-brain intrinsic activity in the hippocampus (Fig. 8A and B) and inferior parietal cortex (Fig. 8A and C) showing a decrease in intrinsic activity 3 months after the SFA-enriched diet, whereas the MUFA group displayed no difference.
levels (17). In this study, MUFAs were sufficient to neutralize the aversive effects of SFAs on measures of insulin sensitivity and secretion.

Further, we showed that body fat accumulation was less in mice fed the MUFA-enriched diet than in those fed the SFA-enriched diet. As both groups of mice were offered diets with the same energy intake, the difference in body fat accumulation between the two diet groups was ascribed to the value of fat quality. A study in rats demonstrated that animals fed safflower oil accumulated less body fat compared with beef tallow–fed rats due to an enhanced thermogenesis and fat oxidation (26), which might together with the increase in locomotor activity contribute to less fat mass in the MUFA-fed mice.

On the basis of our electrocorticogram recordings, we further propose that even a slight increase in SFAs was responsible for the reduction in cortical activity. Radiotelemetric analysis clearly showed that cortical activity was lower in mice fed the SFA-enriched diet compared with MUFA-enriched or chow diet. In this context, it should be noted that an intracerebroventricular insulin application into hyperinsulinemic SFA-mice even lowered brain activity, which might contribute to an impairment in activity and cognitive function as present in subjects with impaired glucose tolerance or even diabetes (27).

Electrocorticogram recordings are widely used to define sleep stages, and a large number of laboratory studies of humans as well as epidemiological data suggest that short sleep duration is associated with an increased risk of developing obesity, impaired glucose tolerance, and insulin sensitivity (28,29). Although most of these studies on food intake, sleep, and metabolic alterations are performed with nonprospective and nonobjective methods, we aimed to identify the impact of fat intake on sleep architecture in mice. Consumption of SFAs decreased wakefulness during the active night phase and increased NREM sleep. This is in agreement with other studies where an association was found for partial sleep loss and disturbance of glucose regulation that involves both reduced β-cell responsiveness and lower insulin sensitivity (30). It is known that the lateral and posterior hypothalamic areas contain neurons specifically active during wakefulness (31), and that the medial prefrontal cortex and basal forebrain are counted among wake-promoting regions (32). Although the effect of SFAs was demonstrated in these brain areas, alterations in REM sleep were only seen in the MUFA-fed animal group. The key brain structure for generating REM sleep is the brainstem and especially the pons and adjacent portions of the midbrain (33). These data suggest that the SFA-enriched diet may specifically act on brain areas controlling wakefulness and NREM sleep, whereas the MUFA-enriched diet preferentially alters mechanisms that act on REM sleep; however, the underlying mechanisms remain to be determined.

We showed that systemic insulin and insulin in the brain alters neuronal activity both in the resting state (13,34) and after stimulation with visual food stimuli in human subjects (35). In particular, the effect of insulin on neuronal activity in humans is impaired by obesity (12), genetic variances (36,37), aging (38), and increased levels of plasma fatty acids as demonstrated by MEG measurements, and

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**FIG. 5.** Different sleep-wake patterns in MUFA- and SFA-fed mice and after intracerebroventricular application of palmitic acid. A–C: Diurnal variations of wakefulness (wake), NREM sleep, and REM sleep in MUFA- (gray circles), SFA- (black circles), or chow-fed (white circles) mice. Data (mean ± SEM of 11–24 animals/diet group) are expressed as minutes per hour for each hour over a 24-h episode (light on 07:00 A.M. to 7:00 P.M.). Statistically significant differences between fat-enriched diet groups and control group are depicted by asterisks (*$P < 0.05$ and **$P < 0.005$).

Impact of intracerebroventricularly injected palmitic acid (225 μmol/L total, black circles) on wakefulness (wake) (D), NREM sleep (E), and REM sleep (F) vs. intracerebroventricularly applied vehicle (white circles) in control mice. Data (mean ± SEM of 13–16 animals) are expressed as described above. Significant differences between intracerebroventricular palmitic acid and vehicle application are presented as follows: **$P < 0.005$ and ***$P < 0.001$. 

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further analysis using fMRI was able to identify specific brain areas related to object processing, memory, and locomotion in lean and obese subjects. By applying fMRI techniques to humans that received either SFAs or MUFAs, we determined that SFA-enriched diet led to a decrease in hippocampal and cortical activity whereas an isocaloric intake of MUFAs showed no such effect. The hippocampus is known to control brain activity in the theta frequency band (39,40). We were able to show that theta activity was correlated to locomotor activity during lifestyle intervention (41), and this was underlined by decreased theta activity in animal models of obesity and physical inactivity (16). Therefore, the human and animal data point in the same direction, where SFAs lead to specific changes in brain areas that are linked to locomotion, memory, and sleep. Of note, both enriched diets contained less polyunsaturated fatty acids compared with chow, and further studies are needed to address this issue in detail.

![FIG. 6. Metabolic consequences of fat-enriched diets in body weight–matched C57BL/6 mice. C57BL/6 mice were weaned on MUFA-enriched (gray), SFA-enriched (black), or chow (white) diet for 8 weeks and body weight–matched animals of all diet groups were selected (n = 4/group). A: Intraperitoneally injected glucose during a GTT was cleared slower and less effectively in SFA-fed mice (black circles) compared with controls (white circles) and MUFA-fed animals (gray circles) (#P < 0.05 to MUFA group). B: Increased fasted blood glucose levels in SFA-fed mice compared with MUFA- (P = 0.05) and chow-fed animals. C: Slightly increased serum insulin concentration during the GTT in MUFA-fed mice (gray circles). D: Increased fasted serum insulin concentrations in the SFA-diet animal group (P = 0.137 to control and MUFA group). E: Increased homeostasis model assessment of insulin resistance (HOMA-IR) in SFA-fed mice (P = 0.1285 to control and P = 0.1137 to MUFA-fed group). F: Analysis of basal power spectral density (indicated for delta [0.5–4 Hz], theta [4–8 Hz], alpha [8–12 Hz], and beta [12–30 Hz] frequencies) using FFT and estimated by ECoG in MUFA- (gray bars), SFA- (black bars), and chow-fed (white bars) animals. Basal ECoG measurements were conducted during a 4-day period (**P < 0.005 and ***P < 0.001 compared with controls; ####P < 0.001 between fat-enriched diet groups). Data are presented as mean ± SEM.](diabetes.diabetesjournals.org)
Together, our data strongly suggested that a MUFA-enriched diet displayed beneficial effects on glucose metabolism and locomotor and cortical activity, as well as sleep behavior even in the presence of obesity. By contrast, a comparable intake of SFAs acted as a mediator of insulin resistance in the brain, with its aversive effects on brain activity and locomotion, and was accompanied by altered sleep-wake architecture.

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No potential conflicts of interest relevant to this article were reported.

T.S. and A.M.H. designed the study, researched data, and wrote the manuscript. C.K. researched data of the human intervention study. S.K. performed and analyzed the fMRI analysis in the human subjects. M.B. performed the fatty acid analysis of edible fats. A.F. researched data of the human intervention study and reviewed and edited the manuscript. H.P. designed the human intervention study and reviewed and edited the manuscript. T.S. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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FIG. 8. Response of the human brain to MUFA- and SFA-enriched diet. A: Color-coded T-value map represents significant (P < 0.05 uncorrected, for display) voxels of decreased intrinsic brain activity 3 months after SFA-enriched yogurt consumption compared with the control group. P < 0.001, whole brain. Plots show change of intrinsic activity in the hippocampus (B) and inferior parietal cortex (C) 3 months after SFA-enriched, MUFA-enriched, and control diet. Only SFA-enriched diet revealed a significant decrease in hippocampal and cortical activity (**P < 0.005 and *P < 0.05). Data are mean ± SEM; n = 8 for each group. (A high-quality color representation of this figure is available in the online issue.)