Dietary protein ingested before and during short photoperiods makes an impact on affect-related behaviours and plasma composition of amino acids in mice

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Abstract
In mammals, short photoperiod is associated with high depression- and anxiety-like behaviours with low levels of the brain serotonin and its precursor tryptophan (Trp). Because the brain Trp levels are regulated by its ratio to large neutral amino acids (Trp:LNAA) in circulation, this study elucidated whether diets of various protein sources that contain different Trp:LNAA affect depression- and anxiety-like behaviours in C57BL/6J mice under short-day conditions (SD). In the control mice on a casein diet, time spent in the central area in the open field test (OFT) was lower in the mice under SD than in those under long-day conditions (LD), indicating that SD exposure induces anxiety-like behaviour. The SD-induced anxiety-like behaviour was countered by an α-lactalbumin diet given under SD. In the mice that were on a gluten diet before transition to SD, the time spent in the central area in the OFT under SD was higher than that in the SD control mice. Alternatively, mice that ingested soya protein before the transition to SD had lower immobility in the forced swim test, a depression-like behaviour, compared with the SD control. Analysis of Trp:LNAA revealed lower Trp:LNAA in the SD control compared with the LD control, which was counteracted by an α-lactalbumin diet under SD. Furthermore, mice on gluten or soya protein diets before transition to SD exhibited high Trp:LNAA levels in plasma under SD. In conclusion, ingestion of specific proteins at different times relative to photoperiodic transition may modulate anxiety- and/or depression-like behaviours, partially through changes in plasma Trp:LNAA.

Key words: α-Lactalbumin: Gluten: Photoperiodism: Seasonal affective disorder: Soya protein

In mammals, photoperiod regulates various physiological functions and behaviours, including breeding, metabolism and stress- and affect-related behaviours(1–6). In humans, seasonal affective disorder (SAD) is characterised by the occurrence of depression, hypersomnia, hyperphagia and carbohydrate craving during specific seasons, usually winter(7). Pathogenesis of SAD is known to involve seasonal fluctuations in the brain serotonergic system. First, serotonin (5-HT) levels in the post-mortem hypothalamic tissue from human subjects were lower in the tissue collected in winter compared with the tissue collected in summer(8). Second, in healthy men, estimated 5-HT in the tissue collected in summer(8). Second, in healthy men, estimated 5-HT in the tissue collected in winter compared with the tissue mortem hypothalamic tissue from human subjects were lower

First, serotonin (5-HT) levels in the post-

In accordance with these mechanisms, carbohydrate craving in the SAD patients, which is a typical symptom of SAD, is hypothesised to be a form of self-medication to compensate for the reduction of 5-HT levels in the brain. In accordance with this hypothesis, SAD patients frequently report an activating effect of carbohydrates(10), and a Trp-deficient diet causes a

Abbreviations: 5-HT, serotonin; FST, forced swim test; LD, long-day conditions; LNAA, large neutral amino acids; OFT, open field test; SAD, seasonal affective disorder; SD, short-day conditions; Trp, tryptophan.

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of SAD, including fat sand rats (1), grass rats (3) and Siberian hamsters (6). As laboratory mouse strains including C57BL/6J cannot produce detectable levels of melatonin, which is an important photoperiodic messenger, they were thought to be an inappropriate model for SAD. Recently, however, we found that C57BL/6J mice under short-day condition (SD) exhibited amplified plasma corticosterone rhythms, whereas those maintained under long-day conditions (LD) exhibited no significant rhythmicity (4). In addition, C57BL/6J mice under SD, compared with mice under LD, showed high immobility in the forced swim test (FST) and a low preference for saccharin, which compared with mice under LD (18 h of light (50 lux) and 6 h of darkness (18L6D)) for at least 1 week before the experiment. The control diet (described below) and tap water were provided ad libitum. All animal experiments were conducted in accordance with the Guidelines for Animal Experiments of the Faculty of Agriculture at Kyushu University, as well as the Law (No. 105) and Notification (No. 6) of the Japanese Government.

Methods

Animals

Male C57BL/6J mice (4 weeks old) were purchased from Japan SLC and randomly housed in plastic cages in groups of three. All mice were housed in boxes placed in a room at 25 ± 1°C with LD (18 h of light (50 lux) and 6 h of darkness (18L6D)) for at least 1 week before the experiment. The control diet (described below) and tap water were provided ad libitum. All animal experiments were conducted in accordance with the Guidelines for Animal Experiments of the Faculty of Agriculture at Kyushu University, as well as the Law (No. 105) and Notification (No. 6) of the Japanese Government.

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Diet

The composition of experimental diets is shown in Table 1. All diets contained 170 g of crude protein/1 kg diet. The percentage of crude protein in casein, gluten, soya protein or ß-lactalbumin powder varies (85, 90, 83 and 94 %, respectively); therefore, the amount of the casein, gluten, soya protein or ß-lactalbumin powder used was 200, 189, 205 and 181 g/kg diet, respectively. The amount was balanced with maize starch. On the basis of the nutrient requirements for rodents, diets with casein and soya proteins were supplemented with ß-cystine. The gluten diet was supplemented with ß-lysine, and the ß-lactalbumin diet was supplemented with ß-arginine. All diets were prepared in our laboratory using ingredients shown in Table 1. All diets were mixed with methylcellulose solution (0.25 %) and dried in an air-dried oven to form a pellet. The casein diet was used as the control diet. Trp and branched-chain amino acid levels and Trp:LNAA in the experimental diets are shown in Table 2.

Experimental design

After acclimation, cages of mice were divided into eight groups (three cages with nine animals per group). Mice were group housed, because our preliminary data revealed that social isolation of C57BL/6J mice masked the effect of photoperiod on immobility in the FST (Y Togo, T Otsuka, M Furuse and S Yasa, unpublished results). The first and second groups were assigned as the SD control and LD control groups, respectively (Fig. 1). SD control mice were exposed to LD for the first 3 weeks and then transferred to SD (6L18D) with the casein diet for the rest of the experimental period. The LD control mice were maintained on a casein diet throughout the experimental period. The third to fifth groups were exposed to LD for 3 weeks with the casein diet, and the lighting condition was then changed to SD (6L18D) with replacement of the casein diet with one of gluten, soya protein or ß-lactalbumin (Fig. 1).

| Table 1. The composition of experimental diets (g/kg diet) |
|----------------------------------|---------------------|---------------------|---------------------|---------------------|
| Protein powder                  | Soya protein        | Gluten              | ß-Lactalbumin       |
| Maize starch                    | 200*                | 189*                | 205*                | 181*                |
| Maize starch                    | 529-486             | 540-486             | 525-866             | 550-586             |
| Sucrose                         | 100                 | 100                 | 100                 | 100                 |
| Soybean oil                     | 70                  | 70                  | 70                  | 70                  |
| Cellulose                       | 50                  | 50                  | 50                  | 50                  |
| Mineral mix†                    | 35                  | 35                  | 35                  | 35                  |
| Vitamin mix†                    | 10                  | 10                  | 10                  | 10                  |
| L-Cys                           | 3                   | 3                   | 1                   | 1                   |
| L-Lys                           | –                   | –                   | 1-6                 | –                   |
| L-Arg                           | –                   | –                   | –                   | 0-9                 |
| Choline bitartrate              | 2-5                 | 2.5                 | 2-5                 | 2.5                 |
| Butylhydroquinone               | 0-014               | 0-014               | 0-014               | 0-014               |

* Crude protein levels are identical to 170 g/kg diet in all diets.
† The vitamin and mineral mixes were purchased from KBT Oriental and contain 97 and 22 % sucrose, respectively.

| Table 2. Tryptophan and related amino acid compositions in protein powders |
|----------------------------------|---------------------|---------------------|---------------------|
| Trp (g/kg)                       | Soya protein        | Gluten              | ß-Lactalbumin       |
| 11                               | 14                  | 7.9                 | 44.3                |
| BCAA (g/kg)                      | 190                 | 168                 | 117.2               | 213                 |
| Trp:LNAA                         | 0.0388*             | 0.0543*             | 0.0405*             | 0.1287*             |

BCAA, branched-chain amino acids; LNAA, large neutral amino acids. * The ratio was calculated on the basis of the amino acid compositions analysed by Japan Food Research Laboratories (casein, gluten and ß-lactalbumin) or Fuji Oil Co., Ltd (soya protein).
These groups were considered the LD-SD protein (SDp) groups. The sixth to eighth groups were exposed to LD for 3 weeks while on a gluten, soya protein or α-lactalbumin diet, and they were then exposed to SD while on a casein diet (Fig. 1). These groups were considered the LD protein (LDp)-SD groups. Food intake (as a cage) in all groups was measured every week, and the total during weeks 2–3 and 5–6 after the start of the experiment was calculated for analysis. After 3 weeks of the last switch of the light/dietary conditions, open field test (OFT) and FST were performed at 2-d intervals. Photoperiod and dietary conditions were continued during these behavioural tests. Mice were euthanised 3, 9 or 21 h after the lights-on (three animals per time point per group) with isoflurane 4 d after the FST, and plasma for analysis of Trp:LNAA and gastrocnemius for muscle fibre type 4–5 d after the FST. Gl, gluten; soya, soya protein; Lac, α-lactalbumin).

**Behavioural tests**

OFT and FST were performed to address the effect of the experimental diets on mood-related behaviours. The tests were performed during light phase under 5 lux light, starting 2 h after the lights-on, because mood-related behaviours and the brain serotonergic system is highly sensitive to photoperiod during light phase in mice (5). Tests in all mice in all groups were completed within 2 h. The OFT was used to analyse spontaneous activity and anxiety-like behaviour in a novel environment. The use of the FST was advocated by a previous study (22), and it is widely used to analyse behavioural despair or depression-like behaviour.

The OFT was performed in the white square field (40×40 cm) with 40-cm high walls. Each mouse was placed in the centre of the apparatus and recorded using digital video for 5 min. Open field behaviour was analysed using ANY-maze software (Stoelting Co.) by dividing the field into 25 squares (5×5 grid). The number of grid lines that were crossed was used as the parameter of spontaneous activity, and the time spent in the central nine grids was used as the parameter of anxiety-like behaviour.

In the FST, mice were individually put into plastic, transparent cylinders (27-cm high, 17-cm diameter) containing 14.5-cm-high water at 25 ± 1°C. The behaviours were recorded using digital video for 7 min, and immobility time during the last 5 min of the 7 min was blindly analysed. The mouse was considered immobile when it floated with its head above the water without moving its tail.

**Measurements of plasma NEFA and amino acids**

Blood samples were obtained by decapitation, and plasma was collected after centrifugation at 3000 g for 10 min at 4°C. The plasma samples were stored at ~80°C until they were analysed. Plasma NEFA levels were measured using the NEFA C-test kit WAKO (Wako Pure Chemical Industries Ltd) according to the manufacturers’ protocol. Free amino acids in the plasma were determined by HPLC (Pico-tag™, Waters) according to a previous method (23). The following was added to the samples: 1 M-sodium hydroxide, calibrated to pH 7.0, and a mixed solution that consisted of 40% 1 M-sodium acetate, 40% methanol and 20% triethylamine. After drying, these samples were allowed to derivatise with a reaction solution that consisted of 70% methanol, 10% triethylamine and 10% phenylisothiocyanate. The samples were dried again, dissolved in 200 µl of Pico-tag sample diluent and filtered.
through a Syringe Driven Filter Unit (Milllex-LG; Millipore) to remove the solid contents. A standard amino acid mixture (Type AN II and type B; Wako Pure Chemical Industries Ltd) was applied using the same methods. These derivatised samples and standards were applied to the HPLC system and equilibrated with buffer A (70 m M-sodium acetate, acetonitrile (975:25)) and eluted with buffer B (water, acetonitrile, methanol (40:45:15)) at a flow rate of 1 ml/min at 46°C.

**SDS-PAGE**

The gastrocnemius was dissected and quickly frozen with liquid N2. Muscle fibre types were analysed by SDS-PAGE of myosin heavy-chain isoforms, IIa, IIb and IIX, according to the previous report (24). Muscle samples were homogenised in a solution containing 10% w/v SDS, 40 mM-dithiothreitol, 5 mM-EDTA and 0.1 M-Tris-HCl buffer. The separating gel consisted of 35% v/v glycerol, 8% w/v acrylamide-N,N'-methylenebisacrylamide (99:1), 0.2 M-Tris-HCl (pH 8.8), 0.1 M glycine, 0.4% w/v SDS, 0.1% w/v ammonium persulphate and 0.05% v/v N,N,N',N'-tetramethylethlenediamine. Samples containing 100 ng of protein were applied to the gel and electrophoresed at a constant voltage of 140 V for 23 h except for the first 40 min, where the maximum current was limited to 10 mA. The gel unit was placed at 4°C throughout the electrophoresis. After the run, gels were stained using a silver stain kit (Silver Stain KANTO III; Kanto Chemicals) and dried using a Gel Dry System (Tefco). These methods have been shown to separate myosin heavy-chain isoforms in a specific manner, as confirmed by Western blotting using antibodies against each isoform (24). We could not clearly separate the bands of IIa and IIX, and thus the bands of IIb and IIA + IIX were quantified densitometrically using the ImageJ 1.47 software (National Institutes of Health). The data are expressed as percentage of IIb content (fast-twitch fibre) to the sum of IIa, IIB and IIX contents.

**Statistical analysis**

LD-SDp and LDp-SD groups were analysed separately. SD and LD controls were used for the control for both the LD-SDp and LDp-SD groups. One-way ANOVA was used to analyse the effects of the diets on food intake, body and epididymal fat weights, plasma NEFA concentrations, OFT, FST and Trp:LNAAs. When significance (P < 0.05) was detected, a post hoc test was performed using Dunnett’s multiple comparison test to compare the values in the SD or LD control group. Behavioural parameters and Trp:LNAAs were also tested using linear mixed models (SAS, PROC MIXED; SAS Institute) with photoperiod or diet as a fixed factor to evaluate the contribution of these factors. In case of Trp LNAAs, time of day was treated as a random effect.

**Results**

**Effects of various proteins on food intake, body weight, epididymal fat mass and plasma NEFA concentrations**

In the LD-SDp groups, food intake during weeks 5–6 but not weeks 2–3 differed significantly (P = 0.0134) between groups. The mice that were on an α-lactalbumin diet showed significantly (P < 0.01) lower levels of food intake compared with those in the SD and LD control groups (Fig. 2a). In the LDp-SDp groups, food intake from weeks 2–3 but not from weeks 5–6 was significantly (P = 0.0032) different between diets (Fig. 2b). Post hoc analysis revealed that mice fed the gluten, soya protein or α-lactalbumin diet ate less food than those in the SD control group (Fig. 2c) and (d)). In the LD-SDp groups, body and epididymal fat weights were higher in the SD control group than in the LD control group (body weight: P < 0.01, fat: P < 0.05). In the LD-SDp group, body and epididymal fat weights significantly differed by diet (body weight: P < 0.0001, fat: P = 0.0012). The mice that were on an α-lactalbumin diet exhibited significantly (body weight: P < 0.0001, fat: P < 0.01) lower levels of body and epididymal fat weights compared with those in the SD control group (Fig. 2c) and (e)). Similar results were obtained in the LDp-SD group (body weight: P = 0.0138, fat: P = 0.0081); both body and epididymal fat weights in the mice that were on an α-lactalbumin diet were significantly (body weight: P < 0.05, fat: P < 0.01) lower than those in the SD control group (Fig. 2d) and (f)). When epididymal fat mass was divided by body weight in each mouse, the mice that were on an α-lactalbumin diet in the LD-SDp groups but not in the LDp-SD groups exhibited significantly (P < 0.05) lower levels compared with those in the SD control group (Fig. 2g) and (h)).

Plasma concentrations of NEFA significantly varied in both LD-SDp (P = 0.0142) and LDp-SD (P = 0.0102) groups (Fig. 2i) and (j)). Post hoc analysis revealed that the NEFA levels were higher in the SD control group than in the LD control group (P < 0.05 in LD-SDp, P < 0.01 in LDp-SD). In the LDp-SD group, mice that were on an α-lactalbumin diet exhibited significantly lower levels of NEFA than those in the SD control group (P < 0.05).

**Effects of various proteins on behaviours in the open field test and forced swim test**

In the OFT, the number of grid lines crossed was not affected by photoperiod or diets with various proteins in both the LD-SDp and LDp-SD groups (Fig. 3a) and (b)). In the LD-SDp groups, the time spent in the centre area is significantly affected by photoperiod (P = 0.0228) and diet (P = 0.0089). One-way ANOVA revealed that it differed significantly between groups (P = 0.0085); the SD control group exhibited significantly (P < 0.05) lower levels than the LD control group (Fig. 3c). The SD-induced suppression of the time spent in the centre area was restored by an α-lactalbumin diet (P < 0.01) (Fig. 3c). Alternatively, in the LDp-SD groups, significant effect of photoperiod (P = 0.0122) and diet (P = 0.0426) was detected. A significant (P = 0.0194) variation was detected among groups, and the reduction in time spent in the centre area under SD was prevented by prior ingestion of a gluten diet (P < 0.05) (Fig. 3d).

In the FST, there was no significant effect of photoperiod or experimental diets on the immobility in the LD-SDp
Fig. 2. Effects of various protein diets (glt, gluten; soya, soya protein; lac, α-lactalbumin) on food intake (a, b), body weight (c, d), fat weight (e, f), fat weight divided by body weight (g, h) and plasma levels of NEFA (i, j) in the long-day condition (LD)-short-day condition (SD) protein (SDp) (a, c, e, g, i) and LD protein (LDp)-SD (b, d, f, h, j) groups. The data of the SD and LD controls were depicted in both the LD-SDp and LDp-SD groups. Values are means (n = 9 except for food intake, which was measured as a cage (three cages in each group)), with their standard errors are represented by vertical bars. Statistically significant difference: * P < 0.05, ** P < 0.01, *** P < 0.001, **** P < 0.0001 (Dunnett’s test) v. SD control: † P < 0.05 (Dunnett’s test) v. LD control.
groups (Fig. 3(e)). In the LDp-SD groups, the experimental diets, but not photoperiod, significantly ($P = 0.0176$) affected immobility. A significant ($P = 0.0248$) variation was detected among groups; the mice that were on a soya protein diet exhibited significantly ($P < 0.01$) lower immobility compared with the SD control group (Fig. 3(f)).

Plasma tryptophan:large neutral amino acids and muscle fibre types

In the LD-SDp groups, the experimental diets, but not photoperiod, significantly ($P = 0.0099$) affected plasma Trp:LNAA. A significant ($P = 0.0248$) variation was detected among groups; post hoc analysis revealed that an $\alpha$-lactalbumin diet under SD significantly increased the ratio to the level in the LD control group (Fig. 4(a)). Trp:LNAA in the LDp-SD groups were affected by diet ($P = 0.0027$) and photoperiod ($P = 0.0265$) with significant variations among groups ($P = 0.0038$). Post hoc analysis revealed that the Trp:LNAA in the SD control group were significantly ($P < 0.05$) lower than those in the LD control group. In contrast to the LD-SDp groups, ingestion of gluten and soya protein diets before exposure to SD significantly (gluten: $P < 0.01$, soya protein: $P < 0.05$) prevented the SD-induced suppression of Trp:LNAA (Fig. 4(b)). When correlation analysis was performed between Trp:LNAA and time spent in the centre area in the OFT, the values in the LDp-SD groups but not those in the LD-SDp groups were significantly correlated (LD-SDp: $P = 0.2469$, $r = 0.1952$, LDp-SD: $P = 0.0174$, $r = 0.3996$) (Fig. 4(c) and (d)). With regard to muscle fibre type, the SD control mice exhibited a higher percentage of IIb isoforms compared with the LD control mice ($P < 0.0001$ in LD-SDp, $P < 0.001$ in LDp-SD) (Fig. 4(e) and (f)). However, in both the LD-SDp and LDp-SD groups, the experimental diets did not affect these ratios (Fig. 4(e) and (f)).
Discussion

In the OFT, SD control mice spent lesser time in the central area compared with LD control mice, which indicated anxiety-like behaviour. This is consistent with the findings of our previous study(5), although the previous study showed photoperiod-induced anxiety-like behaviour using the elevated plus-maze test, but no effect was detected using the OFT. This discrepancy may be because of the differences in photoperiodic conditions; in the present study, LD and SD mice were housed under 18L:6D and 6L:18D, respectively, whereas in the previous study they were housed under 16L:8D and 8L:16D. The larger difference between SD and LD in the present study might enhance the photoperiod-induced anxiety-like behaviour.

The present study further clarified that replacement of casein protein with α-lactalbumin, which has a high Trp:LNAA, during SD significantly increased the time spent in the central area in the OFT (LD-SDp, Fig. 3(c)). This is consistent with the previous report that an α-lactalbumin diet caused an anxiolytic-like effect in OFT or elevated plus-maze test under 12L:12D in C57BL/6J mice and rats(25,26). These changes are unlikely to be a consequence of increased levels of spontaneous activity, because the number of lines crossed did not change with photoperiod. These results indicate that ingestion of α-lactalbumin during seasons with short photoperiods, that is, fall and winter, can improve the anxiety-related problems that occur during these seasons(27,28).

The mechanisms by which α-lactalbumin regulates anxiety-like behaviours remain unclear. A possible explanation is that high Trp:LNAA in the α-lactalbumin diet enhance 5-HT signalling in the brain because of increased transport of plasma Trp through the blood–brain barrier, as plasma Trp:LNAA levels highly correlate with levels of brain Trp (r 0.95), and the sum of 5-HT and its major metabolite 5-hydroxyindoleacetic acid.
In the FST, immobility in SD control mice was slightly lower than that in LD control mice. Ingestion of soya protein before the transition from LD to SD suppressed the immobility during SD compared with the SD control mice (LDp-SD, Fig. 3(f)). Soya protein intake was lower, similar to what was observed with gluten protein intake, and the link between limited energy and subsequent increase of Trp:LNAA may partially explain the results. However, other factors regulating FST behaviour are possible, because the results in the FST were highly different from the results of the OFT, in which α-lactalbumin and gluten in the LDp-SD and LDp-SD groups, respectively, were effective in regulating the time spent in the central area. Anxiety- and depression-like behaviours are controlled by a distinct pathway. 5-HT receptor 2A-deficient mice exhibited more anxiety-like behaviours compared with wild-type mice, whereas no difference was observed in depression-like behaviours. In contrast, the mice deficient in Trp hydroxylase (TPH2), a rate-limiting enzyme in 5-HT synthesis, demonstrated modulation of depression-like, but not anxiety-like, behaviours. These reports suggest that the synthetic pathway of 5-HT is specifically involved in regulation of depression-like behaviours. Although there is evidence that soya phyto-oestrogen induces TPH protein content in the brains of primates, our data cannot be explained by a direct effect of soya protein, given that the immobility in the FST in the mice in LDp-SD was not modified by soya protein diet. Considering that energy intake was limited in the mice fed soya protein in the LDp-SD group, and that insulin sensitivity is enhanced by energy restriction, enhanced action of insulin after the switch to casein might enhance the brain 5-HT signals. In support of this hypothesis, insulin stimulates TPH2 expression in the brains of chickens. Alternately, phyto-oestrogen in soya protein diet under LD might induce long-term effect of oestrogen-related processes involved in regulation of aggression and anxiety, given that sensitivity to oestrogen is regulated by photoperiod in mice, and early exposure to oestradiol alters long-term behavioural sensitivity to anxiolytic drugs in rats.

In our study, body and epididymal fat weights, as well as NEFA levels in plasma, were higher in the SD control group than in the LD control group, whereas food intake was not different between them. Previous studies that used Siberian hamsters or Fischer 344 rats, which are both long-day breeders, reported that body and epididymal fat weights were higher under LD than SD. In contrast, Syrian hamsters show higher body weight under SD than LD. Similarly, in humans, body weight increases from fall to winter. These reports indicate that the direction of the photoperiodic response of body weight and fat mass differs between species. The present study further showed the photoperiodic changes in the ratios of muscle fibre types in C57BL/6j mice; ratio of IIB isoforms, which represent fast-twitch fibres, to the sum of IIA and IIX isoforms, which represent intermediate types of fast- and slow-twitch muscle fibres, are lower under LD than SD. These changes in muscle fibre type might be related to photoperiodic changes in fat metabolism. Type IIA fibres have a higher capacity of oxidative metabolism than type IIB fibres. High IIA/x fibre types in the mice under LD might be related to mobilisation of fatty acids through lipolysis of adipose tissue; released free fatty acids might be transported to muscle for further oxidation. This hypothesis is in line with the report that...
tribbles homolog 3 overexpression, which promotes fatty acid oxidation, causes shift of muscle fibre types from fast- to slow-twitch fibres (50), and can reasonably explain low mass of epididymal fat and plasma NEFA levels observed in our study.

The mice on an α-lactalbumin diet during SD exhibited lower body and epididymal fat weights compared with the SD control mice. This is probably because of the reduced intake of α-lactalbumin diet. This finding is consistent with the Ldp-SD group results in which the mice on an α-lactalbumin diet before the transition to SD exhibited suppressed food intake during the period in which they were fed an α-lactalbumin diet, as well as reduced body and epididymal fat weight, and reduced NEFA levels as a consequence. Notably, the mice in the Ldp-SD group consumed an equivalent amount of the casein diet after the transition to SD, suggesting that the reduction of body and epididymal fat weights was not a consequence of reduced food intake. Our study also showed that composition of muscle fibre type was unmodified by an α-lactalbumin diet, indicating that the effect of this diet on fat and body weights is unrelated to muscle fibre type. Consistent with the findings of our study, several other studies reported that intake of α-lactalbumin resulted in reduction of body and epididymal fat weights in C57BL/6J mice (25, 51).

In conclusion, our study demonstrated that various dietary proteins ingested before and during SD regulate anxiety- and depression-like behaviours, plasma Trp/LNAA, body weight and fat mass. Ingestion of appropriate proteins during specific seasons may be useful to improve or prevent winter-induced mood and metabolism problems, including symptoms of SAD.

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S.Y. designed the study. T. O., R. G., A. I., M. K. and S. S. conducted the animal experiment. Y. O. and W. M. performed the analysis of muscle fibre type. S. Y., T. O. and M. F. analysed the data. S. Y. and T. O. wrote the paper. All authors discussed the results and commented on the manuscript.

The authors declare that they have no conflicts of interest.

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