Effect of tryptophan supplementation on diet-induced non-alcoholic fatty liver disease in mice

Yvonne Ritz, Gyöngyi Bárdos, Astrid Hubert, Maureen Böhle and Stephan C. Bischoff*

Department of Nutritional Medicine, University of Hohenheim, Fruwirthstrasse 12, Stuttgart 70599, Germany

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Abstract

Intestinal serotonin (5-hydroxytryptamine, 5-HT) metabolism is thought to play a role in gut functions by regulating motility, permeability and other functions of the intestine. In the present study, we investigated the effect of tryptophan (TRP), the precursor of 5-HT, supplementation on intestinal barrier functions and non-alcoholic fatty liver disease (NAFLD). An established mouse model of NAFLD induced by feeding a fructose-rich diet (N group) was used in the present study. TRP was administered orally for 8 weeks to C57BL/6j control or NAFLD mice. NAFLD-related liver parameters (hepatic TAG and Oil Red O staining), intestinal barrier parameters (tight-junction protein occludin and portal plasma lipopolysaccharides (LPS)) and 5-HT-related parameters (5-HT, 5-HT transporter (SERT) and motility) were measured. We observed reduced duodenal occludin protein concentrations (P=0.0007), high portal plasma LPS concentrations (P=0.005) and an elevated liver weight:body weight ratio (P=0.01) in the N group compared with the parameters in the control group. TRP supplementation led to an increase in occludin concentrations (P=0.0009) and consecutively reduced liver weight:body weight ratio (P=0.009) as well as overall hepatic fat accumulation in the N group (P=0.05). In addition, the N group exhibited reduced SERT protein expression (P=0.002), which was normalised by TRP supplementation (P=0.02). For the first time, our data indicate that oral TRP supplementation attenuates experimental NAFLD in mice. The underlying mechanisms are not clear, but probably involve stabilisation of the intestinal barrier in the upper small intestine and amelioration of the dysregulated intestinal serotonergic system.

Key words: Tryptophan: Non-alcoholic fatty liver disease: Intestinal permeability: Serotonin: Serotonin re-uptake transporter

The prevalence of non-alcoholic fatty liver disease (NAFLD) is increasing worldwide, with up to 33% of adults being affected in Western countries(1). This alarming increase in the number of individuals affected is related to the contemporary pandemic of obesity fostered by an unhealthy dietary pattern and a sedentary lifestyle, consequently leading to an increase in the risk of death(2). NAFLD patients exhibit signs of liver inflammation and increased hepatic lipid accumulation(3). In addition, the development of NAFLD in obese individuals is closely associated with insulin resistance and other metabolic disorders and thus might be of clinical relevance(4). We and others demonstrated that not only overfeeding but also particular nutrients such as fructose and SFA promote NAFLD in both animals and humans(5–10). Our findings might explain, at least in part, why many, but by far not all, obese individuals develop the metabolic syndrome and associated diseases. In addition, we provided evidence for the involvement of the intestinal serotonergic system in the development of NAFLD and obesity(7,11,12).

Intestinal serotonin (5-hydroxytryptamine, 5-HT) is released by enterochromaffin cells and neurons and is regulated via the serotonin re-uptake transporter (SERT)(15). The SERT is located on epithelial cells and neurons in the intestine(13). Previously, we had tested the hypothesis that the dysregulation of 5-HT due to a diet-induced reduction or a genetically induced loss of the SERT is involved in monosaccharide-induced NAFLD(7). We clearly demonstrated that impairment of SERT promotes the fructose- and glucose-induced damage of the intestinal barrier leading to lipopolysaccharide (LPS) translocation and liver inflammation causing NAFLD(7). Our findings are consistent with those of studies showing that 5-HT is a key regulator not only of intestinal motility(14) but also of inflammation(15,16) and intestinal permeability(17–19). 5-HT is synthesised from the amino acid tryptophan (TRP). Therefore, TRP supplementation is thought to enhance 5-HT synthesis; however, the substrate seems to be a minor regulator of synthesis compared with the rate-determining enzymes tryptophan hydroxylase (TPH1) and TPH2(20). Nevertheless, in earlier studies, TRP supplementation has been shown to result in elevated small-intestinal 5-HT concentrations as well as in increased duodenal and ileal motility(21,22). Concerning the

Abbreviations: 5-HT, serotonin; C group, control diet; C + TRP group, control diet supplemented with tryptophan; LPS, lipopolysaccharide; MZ, mouse breeding; NAFLD, non-alcoholic fatty liver disease; N group, NAFLD mice fed a fructose-rich diet; N + TRP group, NAFLD mice fed a fructose-rich diet supplemented with tryptophan; SERT, serotonin re-uptake transporter; TPH, tryptophan hydroxylase; TRP, tryptophan.

* Corresponding author: Professor Dr S. C. Bischoff, fax +49 711 459 24343, email bischoff.stephan@uni-hohenheim.de
induced fatty liver disease. Supplementation to be a new approach for attenuating diet-development of NAFLD. The present results indicate TRP LPS from the intestine to the liver and the subsequent
teaches is patchy and confusing. Thus, in the present study, we aimed to investigate the effect of TRP supplementation on
the one hand, 5-HT supports liver regeneration after partial hepatectomy and, on the other hand, promotes steato-
hepatitis. Until today, studies concerning TRP supplementation are patchy and confusing. Thus, in the present study,
we aimed to investigate the effect of TRP supplementation on NAFLD. In particular, we were interested in determining
whether TRP supplementation affects the translocation of LPS from the intestine to the liver and the subsequent
development of NAFLD. The present results indicate TRP supplementation to be a new approach for attenuating diet-
induced fatty liver disease.

Materials and methods

Animals and treatments

Mice were housed in a pathogen-free barrier facility accredited by the Association for Assessment and Accreditation of
Laboratory Animal Care. All procedures were approved by the local Institutional Animal Care and Use Committee
(Regional Council Stuttgart). C57BL/6j mice aged 6–8 weeks were obtained from Janvier (Le-Genes-St-isle, France). The
animals were divided into four groups of five to six mice and were fed different diets ad libitum over an 8-week period:
(a) control diet (C group) (water and mouse breeding (MZ) diet; sniff®, containing: 6 % fat, 23 % protein, 5.2 %
sugars, vitamins and minerals); (b) NAFLD-inducing fructose-rich diet (N group) (30 % fructose solution and enriched
MZ diet containing 7.7 % fat, 27.6 % protein, 8.2 % sugars and adjusted vitamins and minerals); (c) control diet supplemented
with TRP (C+TRP group) (0.4 g/kg body weight corresponding to 0.24 g/100 g diet); (d) fructose-rich diet supplemented
with TRP (N+TRP group). The two animal groups fed the fructose-rich diets consumed about 50 % less food (see
Table 1), as has been reported earlier (5), and were therefore
adjusted vitamins and minerals); (b) NAFLD-inducing fructose-rich diet (N group) (30 % fructose solution and enriched
MZ diet containing 7.7 % fat, 27.6 % protein, 8.2 % sugars and adjusted vitamins and minerals); (c) control diet supplemented
with TRP (C + TRP group) (0·4 g/kg body weight corresponding to 0·24 g/100 g diet); (d) fructose-rich diet supplemented
with TRP (N + TRP group). The two animal groups fed the fructose-rich diets consumed about 50 % less food (see
Table 1), as has been reported earlier (5), and were therefore
fed a more energy-dense MZ diet to compensate partially for
the reduced food intake. The amount of TRP administered
was selected according to the method of Rogers & Pesti (24),
who demonstrated reduced hepatic lipid accumulation in
laying hens after TRP supplementation. Liquid and food
intakes, as well as body weight, were recorded weekly.
After 8 weeks of feeding, the mice were anaesthetised with
80mg/kg body weight of ketamine and 6mg/kg body
weight of xylazine, both administered intraperitoneally;
blood was collected from the portal vein just after anaestheti-
sation. After killing the mice, tissue samples of the small
intestine and liver were collected and frozen immediately in
liquid N2 or fixed in paraformaldehyde and frozen in OCT
Tissue-Tek (Sakura Finetek) for further investigation.

Determination of intestinal motility

A charcoal solution (0·5 g charcoal, 0·25 g gum Arabic and
5 ml NaCl (0·9 %)) was orally administered to the mice at a
concentration of 15 μl/kg body weight 10 min before killing
to determine small-intestinal motility. The migration of the
charcoal solution from the pylorus to the small intestine was
measured once the intestine was removed and gently, without
tension, stretched.

Determination of hepatic TAG concentrations

Liver tissue samples weighing 50–100 mg were homogenised
in ice-cold double-concentrated PBS. Tissue lipids were
extracted with methanol (100 μl–chloroform (200 ml), dried
and resuspended in 5 % fat-free bovine serum albumin. Colori-
metric assessment of TAG concentrations was carried out
using a commercially available kit (Randox). Values were
normalised to protein concentrations determined using the
Bradford assay in the liver homogenates (Bio-Rad Laboratories).

Oil Red O and haematoxylin and eosin staining of liver
tissue sections

To determine hepatic lipid accumulation, paraformaldehyde-
fixed liver tissue sections (10 μm) were stained with Oil

<table>
<thead>
<tr>
<th>Table 1. Effect of tryptophan supplementation on physiological parameters§</th>
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<td>(Mean values with their standard errors; n = 5–6)</td>
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<td>Liquid intake (ml/week)</td>
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<td>Food intake (g/week)</td>
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<td>Total energy intake (kJ/week)</td>
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<td>Weight gain (g)</td>
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<td>Liver weight:body weight ratio (%)</td>
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<td>Motility (cm)</td>
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<td>Serotonin content (ng/mg)</td>
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* Mean values were significantly different from those of the control group (P < 0.05).
† Mean values were significantly different from those of the C + TRP group (P < 0.05).
‡ Mean value was significantly different from that of the N + TRP group (P < 0.05).
§ The detailed feeding protocols of the four animal groups are described in the Materials and methods section.
Red O (Sigma) for 10 min, washed and counterstained with haematoxylin (Sigma) for 45 s. Liver histology was assessed using haematoxylin and eosin-stained liver tissue sections. Representative photomicrographs were captured at 400× (Oil Red O) or 200× (haematoxylin and eosin) magnification using a system incorporated in a microscope (Axiovert 200M; Zeiss). The Kleiner et al. (26) scoring system for NASH was applied to two selected mice for each group (26). Selection criteria for the mice were hepatic TAG values that were close to the mean values of each group. The semi-quantitative analysis suggested by Kleiner et al. (26) was used for determining the grade of steatosis (0–3), hepatocyte ballooning (0–2), lobular inflammation (0–2) and fibrosis (0–4). For studying the histological features of NASH, eight haematoxylin and eosin-stained liver tissue pictures per mouse were screened. The hepatic histological feature scores obtained at the onset of NASH are given in Table 2. In addition, the NASH activity score proposed by Kleiner et al. (26), which is the sum of the scores obtained for steatosis, lobular inflammation and hepatocyte ballooning, was determined (Table 2) (26).

Lipopolysaccharide assay
Plasma samples derived from portal vein blood were heated at 75°C for 20 min. The plasma concentrations of LPS were determined using a commercially available limulus amebocyte lysate assay with a concentration range of 0.015–1.2 endotoxin units/ml (Charles River Laboratories International, Inc.).

Determination of duodenal serotonin concentrations
Frozen duodenal tissue samples were homogenised (about 30 s) in ice-cold 0.05 M HCl containing 0.1% ascorbic acid and centrifuged at 14,000 rpm for 5 min at 4°C. The supernatant was immediately acylated (37°C for 15 min in a water-bath) and 5-HT concentrations were determined within 1 h to ensure stability. The concentrations of 5-HT (expressed in ng/mg tissue weight) in the supernatant were determined at room temperature (18–25°C) using an enzyme immunoassay kit (sensitivity of 2.68 ng/ml) according to the manufacturer’s instructions (IBL).

Table 2. Kleiner scoring system* for non-alcoholic fatty liver disease (NASH; n=2)  

<table>
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<tr>
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<th>Steatosis (0–3)</th>
<th>Ballooning (0–2)</th>
<th>Inflammation (0–2)</th>
<th>Fibrosis (0–4)</th>
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<tr>
<td>C</td>
<td>0</td>
<td>0</td>
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<td>C + TRP</td>
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<td>N</td>
<td>1</td>
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<td>N + TRP</td>
<td>0</td>
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C, control diet group; C + TRP, control diet group with tryptophan supplementation; N, NASH-inducing fructose-rich diet group; N + TRP, NASH-inducing fructose-rich diet group with tryptophan supplementation.

*The Kleiner et al. (26) scoring system was used to analyse the development of NASH in haematoxylin and eosin-stained liver tissue sections. The NASH activity score for the N group was 3 (sum of the scores of steatosis, ballooning and inflammation), whereas the score was 0 for all other groups, meaning no histological NASH features were detected.

Immunoblotting
To prepare total tissue protein lysates, snap-frozen small intestine samples were homogenised in a lysis buffer (20 mM 3-morpholin-4-yl propane-1-sulfonic acid (MOPS), 150 mM NaCl, 1 mM EDTA, 1% Nonidet P-40, 1% sodium deoxycholate and 0.1% SDS) containing a protease inhibitor mix (Roche). Protein lysates (30 μg protein/well) were separated on a 10% SDS-polyacrylamide gel and transferred onto Hybond™-P polyvinylidene difluoride membranes (GE). The resulting blots were then blocked for 45 min at room temperature with 1% skimmed milk for the detection of SERT. 5% skimmed milk for the detection of occludin or 2.5% bovine serum albumin solution for the detection of β-actin. After blocking, the blots were incubated overnight at 4°C with primary antibodies against SERT (1:500; Santa Cruz Biotechnology) or occludin (1:500; Zymed Laboratories) and β-actin (1:750; Cell Signaling, New England Biolabs GmbH) as the reference protein. The blots were washed and incubated for 45 min at room temperature with horseradish peroxidase-conjugated secondary antibodies (all from Cell Signaling). The bands were visualised using the SuperSignal Western Dura Kit (Pierce, Thermo Fischer). To ensure equal loading, all the blots were stained with Ponceau red (Sigma-Alrich), and the signals were normalised to β-actin signals. Protein bands were densitometrically analysed using the software AlphaEaseFC (FluorChem™) from Alpha Innotech.

RNA isolation and real-time RT-PCR
Total RNA was extracted from duodenal tissue samples using the TriFast™ reagent (PEQLAB). RNA concentrations were determined spectrophotometrically, and 0.25 μg of total RNA was reverse-transcribed using the iScript 8 DNA synthesis Kit (Bio-Rad Laboratories) followed by a DNase digestion step (Fermentas). PCR primers were designed using the Primer3 software (TPH: forward CATCACGCGAGAACAGTTGA; reverse TTCGGATCCGATCAAGCGA; 18S: forward ATCAGATCCGTGTAGTC; reverse CCAGAGTGCTGTTGAT; Whitehead Institute for Biomedical Research). iScript EvaGreen Supermix (Bio-Rad Laboratories) was used to prepare the PCR mix. The amplification reactions were carried out in an iCycler (Bio-Rad Laboratories) with forty cycles of a two-step PCR (denaturation at 95°C for 35 s, denaturation at 95°C for 5 s, and annealing/extension at 62°C for 10 s). The fluorescence intensity of each sample was measured at each temperature change to monitor the amplification of the target gene. The comparative threshold cycle (CT) method was used to determine the amount of target gene, normalised to that of an endogenous reference gene (18S) and relative to a calibrator (2−ΔΔCt). The purity of PCR products was verified by melting curve analysis and gel electrophoresis.

Statistical analysis
Results are reported as means, with their standard errors. ANOVA or Kruskal–Wallis test was used if variances calculated with Bartlett’s test varied significantly. If there were significant
differences between the groups, we used as a *post hoc* test an unpaired, two-tailed *t* test or *t* test with Welch’s correction (if variances differed significantly), as we were interested in comparing the C and N groups as well as the N and N+TRP groups. A *P* value <0.05 was determined as the level of significance before the start of the study. For calculation and graph design, the software GraphPad Prism 5 (GraphPad Software) was used.

**Results**

*Effect of tryptophan supplementation on physiological parameters*

Liquid intake and weight gain were similar in the four dietary groups (Table 1). Independent of TRP supplementation, we observed, as has been shown previously(5), a reduction in solid food intake by 48.5–51.0% in mice given drinking-water containing fructose (fructose-rich diet) compared with the food intake in mice given plain drinking-water.

*Effect of tryptophan supplementation on hepatic lipid accumulation*

The liver weight:body weight ratio (*P*= 0.014) was significantly increased in the N group (Fig. 1(a)). TRP supplementation reduced the liver weight:body weight ratio to normal levels (*P*= 0.009; Fig. 1(a)). As has been shown previously(5), here fructose-rich diet-fed mice exhibited a dramatic increase in overall hepatic lipid accumulation compared with the fructose-rich diet-fed mice in which hepatic lipid accumulation was reduced by TRP supplementation (Fig. 1(b) and (d)). Similarly, fructose-rich diet-fed mice exhibited a higher degree of steatosis, cellular ballooning and lobular inflammation compared with mice fed the fructose-rich diet with TRP supplementation (Fig. 1(e); Table 2). The NAFLD activity score was 5 for the N group and 0 for all the other groups (Table 2). Nevertheless, hepatic TAG concentrations were not significantly reduced in the TRP-supplemented N group (Fig. 1(c)).

*Effect of tryptophan supplementation on tight-junction protein expression*

We measured the expression of occludin, a tight-junction protein thought to be important for the epithelial barrier function in the small intestine(27). Measurements in the duodenum, the first site of sugar absorption, revealed that occludin protein expression in the N group was significantly reduced (*P*= 0.0007) compared with that in the control group (Fig. 2(a)). Interestingly, TRP supplementation normalised duodenal occludin protein expression in the N group (*P*= 0.0009; Fig. 2(a)). By contrast, in the ileum, despite a tendency for occludin protein expression to decrease in the N and N+TRP groups, TRP supplementation seemed to have no influence (Fig. 2(b)). The concentrations of portal plasma LPS were almost 7-fold increased in the N group compared with that in the control group (*P*= 0.005; Fig. 2(c)). However, TRP supplementation led to no significant reduction in LPS concentrations in the N group (Fig. 2(c)).
diet (Fig. 3). As expected, TPH1 concentrations in the duodenum (Table 1). 

influence neither on motility nor on overall 5-HT concentration by TRP supplementation in addition to fructose-rich diet feeding (data not shown). TRP supplementation had an influence neither on motility nor on overall 5-HT concentrations in the duodenum (Table 1).

Effect of tryptophan supplementation on the intestinal serotonergic system

To determine whether measurable alterations in the intestinal serotonergic system were associated with the protective effect of TRP, SERT protein concentrations, TPH1 mRNA expression and 5-HT concentrations were quantified in the duodenum. Confirming our own previous data(7), we observed a significant reduction in SERT protein concentrations by 56·4% in the N group compared with that in the control group (P=0·002, Fig. 3). TRP supplementation increased SERT protein (P=0·02) expression in mice fed the fructose-rich diet (Fig. 3). As expected, TPH1 mRNA expression was increased (P=0·01) in the N group, but it was not normalised by TRP supplementation in addition to fructose-rich diet feeding (data not shown). TRP supplementation had an influence neither on motility nor on overall 5-HT concentrations in the duodenum (Table 1).

Discussion

There is now growing experimental evidence that alterations in the intestinal barrier resulting in an increased translocation of bacterial products contribute to the development of NAFLD(7,27,28). However, the triggers causing barrier impairment, LPS influx into the liver and hepatic lipid accumulation are largely unknown. We and others have validated that particular dietetic factors such as sugars, in particular, fructose(5), and/or SFA(8,9) play a role, but the underlying mechanisms have been described only rudimentarily.

Recent findings reported by our group have suggested that the intestinal serotonergic system, and especially SERT and 5-HT assessed in the present study, might be involved in the regulation of intestinal permeability and thus translocation of gut-derived bacterial LPS(7). Such data raised the idea that the intestinal serotonergic system could be an interesting pharmacological target for the prevention of fatty liver disease. Therefore, we chose the 5-HT precursor TRP as a dietary supplement, which has been investigated in various animal models(22,23) and has no major side effects on liquid or food intake or on weight gain(29). In the present study, we observed an increase in energy intake after TRP supplementation in the control mice but not in the NAFLD mice with TRP supplementation. The increase in energy intake did not have an effect on weight gain, which might be, among others, due to a slight but not significantly increased motility in the TRP-treated control mice.

In the present study, we clearly demonstrated the protective effect of TRP supplementation on the stabilisation of the intestinal barrier through the enhancement of occludin expression and the obvious reduction of fatty acid accumulation in the liver. The protective effect of TRP supplementation in our NAFLD model was not associated with an attenuation of intestinal translocation of bacterial LPS, possibly because of the partial normalisation of the tight junction impairment only in the duodenum but not in the ileum. Interestingly, L-TRP administration in the rat ileum has been found to reduce net Na+ and fluid absorption, supposedly due to morphological changes(30).

Effect of tryptophan supplementation on occludin protein expression in the duodenum. Occludin protein expression was quantified by Western blotting in the duodenum (a) and ileum (b). Representative blot images of occludin and β-actin as well as the quantitative analysis results of the blots are shown. Values are means, with their standard errors represented by vertical bars (n 4–6). C, control diet; C + TRP, control diet with TRP supplementation; N, non-alcoholic fatty liver disease (NAFLD) mice fed a fructose-rich diet; N + TRP, NAFLD mice fed a fructose-rich diet with TRP supplementation. Mean value was significantly different from that of the C group: ** P<0·01, *** P<0·001. ††† Mean value was significantly different from that of the N group (P<0·001).

Fig. 2. Effect of tryptophan (TRP) supplementation on occludin protein expression in the duodenum. Occludin protein expression was quantified by Western blotting in the duodenum (a) and ileum (b). Representative blot images of occludin and β-actin as well as quantitative analysis results of the blots are shown (a) and (b)). Lipopolysaccharide (LPS; endotoxin units (EU)/ml) concentrations were measured in portal vein plasma (c). Values are means, with their standard errors represented by vertical bars (n 4–6). C, control diet; C + TRP, control diet with TRP supplementation; N, non-alcoholic fatty liver disease (NAFLD) mice fed a fructose-rich diet; N + TRP, NAFLD mice fed a fructose-rich diet with TRP supplementation. Mean value was significantly different from that of the C group: ** P<0·01, *** P<0·001. ††† Mean value was significantly different from that of the N group (P<0·001).

Fig. 3. Effect of tryptophan (TRP) supplementation on duodenal serotonin re-uptake transporter (SERT) protein expression. SERT protein expression in the duodenum was measured by Western blotting. Representative blot images of SERT and β-actin as well as the quantitative analysis results of the blots are shown. Values are means, with their standard errors represented by vertical bars (n 4–6). C, control diet; C + TRP, control diet with TRP supplementation; N, non-alcoholic fatty liver disease (NAFLD) mice fed a fructose-rich diet; N + TRP, NAFLD mice fed a fructose-rich diet with TRP supplementation. ** Mean value was significantly different from that of the C group (P<0·01). † Mean value was significantly different from that of the N group (P<0·05).
TRP supplementation has been found to attenuate the development of NAFLD in laying hens\(^{25,24}\).

We not only found that TRP supplementation attenuated NAFLD but also found that it increased the lower intestinal expression of SERT in the N group. Lower intestinal expression of SERT in our NAFLD model was accompanied by liver steatosis, increased portal plasma LPS concentrations and tight-junction protein loss in the small intestine, as has been postulated here and previously\(^7\). Similarly, SERT knock-out mice have been found to develop liver damage on being fed a glucose-rich diet, which induced no or minor damages in C57BL/6J wild-type mice\(^7\). Our data indicated that fructose is more harmful than glucose in this respect, but in animals lacking SERT, glucose becomes equally harmful with regard to fatty liver disease as fructose in wild-type mice. Therefore, we suggest that SERT and thus the serotoninergic system play a role in sugar-induced liver damage. Possibly, loss of SERT in the intestine leads to a dysregulation of 5-HT signalling that may evoke an enhanced inflammatory response as determined in IL-10 knockout and SERT/IL-10 double-knockout mice\(^{15}\). The latter were characterised by high concentrations of TNF-\(\alpha\), which modulates tight-junction proteins via myosin light-chain kinase activation\(^{53}\). Therefore, we assume that the induction of SERT might be an important mechanism, besides occludin enhancement, with regard to how TRP might attenuate NAFLD at the intestinal level (Fig. 4).

In contrast to previous findings reported by Ozer et al.\(^{21,22}\) no significant differences in small-intestinal 5-HT concentrations or motility were detected among the dietary groups. Ozer et al.\(^{21,22}\) reported 5-HT concentrations and duodenal and ileal motility to be significantly increased in mice treated with TRP (0·1 g/kg body weight per 24 h, intraperitoneally), which might have resulted from the different doses and routes of administration (oral vs. intraperitoneal). However, it is unclear whether overall concentrations of 5-HT in tissues are a reliable marker for 5-HT signalling. Therefore, we determined the mRNA concentration of \(TPH1\), the rate-limiting enzyme of 5-HT synthesis, which was significantly increased on feeding the fructose-rich diet but not normalised by TRP supplementation. Nevertheless, we assume that \(TPH1\) might be a more relevant marker for bioavailable 5-HT at the intracellular level than the overall 5-HT tissue concentration. Of course, other mechanisms such as central effects of TRP cannot be excluded and could contribute to the beneficial effects of TRP in the experimental NAFLD model used in the present study.

Our data provide the first evidence that oral supplementation of the amino acid TRP attenuates NAFLD in mice. In conclusion, we report that modulation of the upper intestinal barrier and the intestinal serotoninergic system via TRP supplementation might be an important mechanism of protection from the development of NAFLD in mice. Adverse effects of TRP supplementation were not observed in our mouse model; however, these have to be considered when testing this approach in human subjects.

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Tryptophan and fatty liver disease

References


