**Ganoderma lucidum** promotes sleep through a gut microbiota-dependent and serotonin-involved pathway in mice

Chunyan Yao1,5, Zhiyuan Wang2,5, Huiyong Jiang3,5, Ren Yan3,5, Qianfei Huang1, Yin Wang1, Hui Xie2, Ying Zou4, Ying Yu1,2,5 & Longxian Lv3,5

_Ganoderma lucidum_ is a medicinal mushroom used in traditional Chinese medicine with putative tranquilizing effects. However, the component of _G. lucidum_ that promotes sleep has not been clearly identified. Here, the effect and mechanism of the acidic part of the alcohol extract of _G. lucidum_ mycelia (GLAA) on sleep were studied in mice. Administration of 25, 50 and 100 mg/kg GLAA for 28 days promoted sleep in pentobarbital-treated mice by shortening sleep latency and prolonging sleeping time. GLAA administration increased the levels of the sleep-promoting neurotransmitter 5-hydroxytryptamine and the _Tph2, Iptr3_ and _Gng13_ transcripts in the sleep-regulating serotonergic synapse pathway in the hypothalamus during this process. Moreover, GLAA administration reduced lipopolysaccharide and raised peptidoglycan levels in serum. GLAA-enriched gut bacteria and metabolites, including _Bifidobacterium_, _Bifidobacterium animalis_, indole-3-carboxylic acid and acetylphosphate were negatively correlated with sleep latency and positively correlated with sleeping time and the hypothalamus 5-hydroxytryptamine concentration. Both the GLAA sleep promotion effect and the altered faecal metabolites correlated with sleep behaviours disappeared after gut microbiota depletion with antibiotics. Our results showed that GLAA promotes sleep through a gut microbiota-dependent and serotonin-associated pathway in mice.

Sleep is important for all living species, and it comprises roughly one-third of our lives. Good sleep is a prerequisite for physical function (including promoting growth, learning and cognitive development), immunity, and poor or insufficient sleep is currently an important health problem. Currently, 15–35% of adults suffer from regular sleep disruptions, such as difficulties in initiating sleep, insufficient sleep time or frequent waking during the night. Sleep problems may damage daytime lives by feeling exhausted and making troubles. Moreover, persistent sleep problems are frequently associated with cardiovascular diseases, obesity, diabetes, and mortality.

_Ganoderma lucidum_, also known as Lingzhi in China, is a medicinal mushroom that has been widely used for medicinal purposes over the past 2000 years, including for tranquilizing. The tranquilizing effect of _G. lucidum_ was recorded in "Shennong's herbal classic" as early as the first century BC and currently in the Chinese Pharmacopoeia. Recently, it was reported that _G. lucidum_ improved the sleep of patients with insomnia or other mental disorders. Furthermore, _G. lucidum_ promoted sleep in pentobarbital-treated mice and rats. However, most sleep-related experiments are based on the water extract of _G. lucidum_. The effect of other extract parts on sleep has not been explored to date.

Recent studies indicate that alterations in the gut microbiota might be associated with sleep through the gut–brain axis. Structural components of the microbial cell wall, such as lipopolysaccharide (LPS) and

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1. Key Laboratory of Nutrition of Zhejiang Province, Institute of Health Food, Hangzhou Medical College, Hangzhou, Zhejiang, China.
2. Animal Center, Hangzhou Medical College, Hangzhou, Zhejiang, China.
3. State Key Laboratory for Diagnosis and Treatment of Infectious Diseases, Collaborative Innovation Center for Diagnosis and Treatment of Infectious Diseases, The First Affiliated Hospital, College of Medicine, Zhejiang University, Hangzhou, Zhejiang, China.
4. The Second Affiliated Hospital of Zhejiang, Chinese Medical University, Hangzhou, Zhejiang, China.
5. These authors contributed equally: Chunyan Yao, Zhiyuan Wang, Huiyong Jiang and Ren Yan.

**5**. E-mail: yuying@hmc.edu.cn; lvlongxian@zju.edu.cn
peptidoglycan (PG), stimulate the innate immune system that begins at the intestinal mucosal surface and impacts the entire body\textsuperscript{14}. Anderson et al. found that better sleep quality was associated with higher proportions of the gut microbial phyla \textit{Verrucomicrobia} and \textit{Lentisphaerae} in healthy adults, suggesting a possible relationship between sleep quality and the gut microbiota\textsuperscript{15}. Poroyko et al. showed that chronic sleep fragmentation (SF) was related to the gut microbiota through conventionalization of germ-free mice with the gut microbiota of mice with SF\textsuperscript{16}.

In this work, the effect of the acidic part of the alcohol extract of \textit{G. lucidum} mycelia (GLAA) on sleep was investigated using a hypnotized mouse model induced by pentobarbital sodium. Furthermore, the underlying mechanism was explored by integrating microbiomics, transcriptomics, and metabolomics and then verified using gnotobiotic mice.

Results

GLAA has no obvious toxicological effect on mice. Body weight was evaluated every week, and administration of GLAA did not affect mouse weight (Fig. 1a). After the mice were sacrificed, the spleen, liver, kidney and thymus were observed by a pathologist who was blind to our experiment, and no obvious pathological changes were found in these organs. Moreover, pre-treatment with GLAA also did not alter the organ/body weight ratio of these organs (Fig. 1b). In addition, administration of GLAA did not induce histological abnormalities in the colons of mice (Supplementary Fig. S1). These results showed that GLAA was safe for mice at dose from 25 to 100 mg/kg.

GLAA potentiates pentobarbital-induced sleep. Using a mouse model of pentobarbital-induced hypnosis, we observed that gavage administration of GLAA for 4 weeks led to significant decreases in sleep latency and increases in sleeping time compared with the solvent control containing 0.05% sodium carboxymethyl acid (0.05% CMC-Na). The average sleep latency of the control group was approximately 300 s, while those of groups administered GLAA at levels ranging from 25 to 100 mg/kg were shortened to approximately 250 s (Fig. 1c). The sleeping time of mice gavaged with 25 mg/kg, 50 mg/kg and 100 mg/kg GLAA increased significantly from 1570.4 s (at 0 mg/kg) to 2816.5 s, 3000 s and 3523.5 s, respectively, showing a certain dose dependency (Fig. 1d). These results suggested that gavage administration of GLAA could significantly promote the sleep quality of mice.

GLAA increases the 5-HT content in the hypothalamus. To determine which neurotransmitters were responsible for the sleep changes in mice, 5-HT, GABA, norepinephrine and dopamine were detected in...
the hypothalamus of mice with a commercial ELISA kit. Only the 5-HT level was significantly increased after administration of 50 mg/kg and 100 mg/kg GLAA (Fig. 1e), indicating that 5-HT may act as a target neurotransmitter for GLAA to promote sleep health in mice.

**GLAA reduces the LPS content and increases the PG content in serum.** LPS and PG are two major components of the bacterial cell wall and two indicators of body inflammation. LPS exists in the cell wall of only gram-negative bacteria, and the PG content in the cell wall of gram-positive bacteria is much higher than that in the cell wall of gram-negative bacteria. In our study, administration of GLAA significantly reduced the LPS content but raised the PG content in serum (Fig. 1f), indicating that GLAA may change the composition of the gut microbiota.

**GLAA changes the composition of the gut microbiota.** Next-generation sequencing of 16S rRNA V3–V4 regions was conducted to investigate the faecal microbiota composition. As shown by the Chao1 index, the microbiota of the control and 100 mg/kg GLAA groups exhibited the same richness (Fig. 2a); as shown by the Simpson index, there was also no significant difference in microbiota diversity between the control group and the 100 mg/kg GLAA group (Fig. 2a). However, the principal coordinate analysis (PCoA) results showed that the microbiota profiles of the two groups were clearly separated from each other (Fig. 2b). PERMANOVA further confirmed that the compositions were different between the control and 100 mg/kg GLAA groups (p = 0.001).

To analyse the effect of 100 mg/kg GLAA on the intestinal microbiota, discriminative taxa were identified by LEfSe analysis at multiple levels, and the most different results (linear discriminant analysis (LDA) score > 3.5) were retained to investigate the alteration in the microbiota structure (Fig. 2c). Compared with the control group, the 100 mg/kg group exhibited enrichment of the family Bifidobacteriaceae and its affiliated genus Bifidobacterium and species Bifidobacterium animalis and species Lactobacillus and species Lactobacillus reuteri, the family Porphyromonadaceae and its affiliated genus Odoribacter, the family Erysipelotrichales and its affiliated genus Turicibacter, and the genus Rikenella. In addition, administration of 100 mg/kg GLAA depleted the family Ruminococcaceae and its affiliated genus Oscillibacter and the genera Anaerotignum and Roseburia. Finally, 100 mg/kg GLAA significantly enriched Bifidobacterium, Rikenella, Odoribacter and Turicibacter at the genus level, as well as Bifidobacterium animalis at the species level after p value correction (q = 0.05, Fig. 2d).

The correlation of discriminative bacterial genera with the sleep behavioural index and concentration of 5-HT in the hypothalamus were determined using Spearman’s rank correlation analysis (Fig. 2e). Bifidobacterium, Odoribacter, Rikenella and Turicibacter at the genus level, as well as Bifidobacterium animalis at the species level, were negatively correlated with sleep latency. Bifidobacterium and Odoribacter at the genus level, as well as Bifidobacterium animalis at the species level, were positively correlated with sleeping time and the concentration of 5-HT in the hypothalamus. These results indicated that the gut microbiota may be involved in the process by which GLAA promotes sleep in mice.

**GLAA alters gut metabolites.** To explore the relationship between alterations in gut metabolites and sleep, faecal samples were submitted to metabolome analysis using LC–MS/MS. In total, 13,023 metabolites were identified. 1037 metabolites with score > 0.5 and a level 2 identification were retained (supplemental material 1). Principal component analysis (PCA) showed that the control group and 100 mg/kg group clustered separately from each other in both the original identification metabolome profiles and the retained metabolome profiles, suggesting that the metabolome profiles of these two groups (Fig. 3a, Supplementary Fig. S2a) are quite different. Similar results were obtained from the orthogonal partial least squares discriminant analysis (OPLS-DA) (Fig. 3b, Supplementary Fig. S2b).

The top 30 metabolites were selected based on the variable importance in the projection (VIP) value from the retained metabolome profiles to explain the difference between the two groups in the OPLS-DA model (Fig. 3c), which most contributed to the alteration of the metabolome between the control group and the 100 mg/kg group. In addition, all the top 30 VIP metabolites showed significant difference (p < 0.05). After gavage administration of 100 mg/kg GLAA, hydroxypropionic acid, malonic acid, dinitramine, esprocarb, ostruthin, d-ribose, (7S,8S)-DiHODE, avermectin B1b aglycone, steviol, d-tagatose, γ-l-glutamyl-l-2-aminobutyrate, His-Phe-Met, 5-androstenetriol, His-Glu-Phe-Glu, soyasaponin II, d-(+)-raffinose, Leu-Asp-Ser, 17α,20α-dihydroxycholesterol and Leu-Arg-Ser-Lys were depleted in the gut.

Spearman’s rank correlation analysis showed that most metabolites were correlated with the sleep behavioural indexes and concentration of 5-HT in the hypothalamus (Fig. 3c). d-ribose and daidzein showed negative correlation with sleep-promotion index and hypothalamic 5-HT concentration. On the contrary, galactosylsphingosine, 2-hydroxy-2-ethylsuccinic acid, Leu-Arg-Ser-Lys, indole-3-carboxylic acid, 17α,20α-dihydroxycholesterol and Leu-Arg-Ser-Lys were correlated with at least one sleep behavioural index, which suggested that they were positively correlated with sleep-promotion effect. Conversely, Cys-Arg-Phe, acetylphosphate, Leu-Arg-Ser-Lys and betaine exhibited negatively correlated with sleep-promotion effect.
Antibiotic-induced depletion of the gut microbiota inhibits the sleep-promoting effect and the increase in 5-HT caused by GLAA. To explore whether the sleep promotion effect of GLAA was dependent on the gut microbiota, we depleted the gut microbiota in mice using antibiotics according to a previous report. After 12 days of depletion of the gut microbiota using an antibiotic concoction, GLAA was gavaged from the 13th day to the 40th day, together with the antibiotic concoction. Notably, neither antibiotic treatment nor GLAA gavage had a significant effect on the body weight of mice (Fig. 4b).

Then, a mouse model of the pentobarbital hypnosis test was established. All the control mice fell asleep within 600 s, while all mice administered 100 mg/kg GLAA fell asleep within 300 s. However, mice pre-treated with antibiotic concoction exhibited only 40% sleep incidence during the experimental period. Even after administration of 100 mg/kg GLAA, the incidence of sleep in mice pre-treated with antibiotic concoction was only 50% (Fig. 4a).

Furthermore, pre-treatment with antibiotic concoction inhibited the increase in 5-HT induced by 100 mg/kg GLAA, as observed in mice (Fig. 4a). Therefore, depletion of the gut microbiota prevented the sleep promotion effect of GLAA, indicating that gut microbiota might play a pivotal role in GLAA-mediated promotion of sleep.

Our results showed that antibiotic treatment depleted more than 85% of the operational taxonomic units (OTUs) of gut bacteria, with a significant decrease in the Chao1 and Simpson indices (Fig. 4c); moreover, the significant decrease in the LPS content in serum further indicated a decrease in microbial quantity in the gut (Fig. 4c). The remaining bacteria mainly included Proteobacteria (99.06%) at the family level (Supplementary Fig. S3a), and Klebsiella (81.4%) and Proteus (13.8%) at the genus level (Supplementary Fig. S3b). In contrast, in the control and 100 mg/kg groups without antibiotics, the microbiota presented a normal distribution of Firmicutes (63.6% and 56.96%), Bacteroidetes (23.5% and 32.38%), Proteobacteria (9.6% and 6.47%), Deferrribacter (2.9% and 2.3%), and Actinobacteria (0.1% and 1.64%), with other phyla accounting for less than 0.2% (Fig. 4d). The above-mentioned microbial markers altered by GLAA were all depleted after antibiotic treatment, and this depletion was not reversed by GLAA in the antibiotic-gavaged mice (Fig. 4e).

Among the top 30 faecal metabolites altered by 100 mg/kg GLAA, the alterations in 13 metabolites were totally or partially prevented by depletion of the gut microbiota (Fig. 4f). Depletion of the gut microbiota prevented the GLAA-induced elevation of acetylphosphate, 2-hydroxy-7-hydroxymethylchromene-2-carboxylate, Pro-Val-Val, Leu-Arg-Ser-Lys and indole-3-carboxylic acid. It also prevented the GLAA-induced depletion of hydroxyproprionic acid, Asp-Ile-Glu and daidzein. Notably, except for esprocarb, the remaining 12 of the above-mentioned 13 metabolites were strongly correlated with sleep latency, sleeping time and/or the concentration of 5-HT in the hypothalamus.

GLAA regulates the serotonergic synapse pathway. Genome-wide transcriptional profiling was performed to analyse the changes in gene transcription in the hypothalamus of mice. In total, 830,782,772 valid sequencing reads were generated from 21 samples (5–6 samples per group), and 55,450 expressed genes were obtained after p value correction. A total of 13 pathways, namely, nitrogen metabolism; d-arginine and d-ornithine metabolism; protein digestion and absorption; prion diseases; NOD-like receptor signalling pathway; glycine, serine and threonine metabolism;...
cytokine–cytokine receptor interaction; alcoholism; glutamatergic synapse; and serotonergic synapse (Fig. 5d). These pathways were mostly involved in sleep (glutamatergic synapse and serotonergic synapse) and immunity (prion diseases, NOD-like receptor signalling pathway and cytokine–cytokine receptor interaction) regulation23-24.

Then, we specifically examined the serotonergic synapse, the output of which is 5-HT. After gavage administration of 100 mg/kg GLAA, the transcription levels of Tph2, Itpr2 and Gng13 in the serotonergic synapse pathway increased significantly, and depletion of the gut microbiota also blocked these increases. Tryptophan is converted by the enzyme tryptophan hydroxylase (TPH) to 5-hydroxytryptophan (5-HTP), and 5-HTP is further converted to 5-HT in presynaptic cells25. 5-HT is secreted into postsynaptic cells and binds to 5-HT receptors (mainly 5-HT1R, 5-HT2R and 5-HT5R) based on the KEGG pathway, thus activating downstream signalling pathway-related genes (including Itpr2 and Gng13) and promoting sleep (Fig. 5e,f).

Figure 3. Gavage administration of GLAA led to alterations in gut metabolites that were correlated with sleep behaviour in mice. (a) PCA score plots of metabolome profiles of the control group and the 100 mg/kg GLAA group. (b) OPLS-DA score plots of metabolome profiles of the control group and the 100 mg/kg GLAA group. (c) Correlations of the top 30 VIP (variable importance in projection) gut metabolites, sleep behaviour and the concentration of 5-HT in the hypothalamus were determined using Spearman’s rank correlation analysis. The image is generated by SIMCA 14.1 and R package “corrplot”. The results with p values < 0.05 are shown in the heatmap. The colour key and circle size indicate the strength of the correlation (r value). Red indicates positive correlations; blue indicates negative correlations. The data are calculated from 8 samples per group. Except specifically mentioned, the images are generated by SIMCA 14.1. GLAA the acidic part of the alcohol extract of G. lucidum mycelia, PCA principal component analysis, OPLS-DA orthogonal partial least squares discriminant analysis, VIP variable importance in projection, 5-HT 5-hydroxytryptamine.
Discussion

Sleep problems have become a hot topic of global concern. *G. lucidum* has been used as a tranquilizing agent for the treatment of restlessness, insomnia, and palpitation in China for hundreds of years. Chu et al. reported that 80 mg/kg and 120 mg/kg water extracts of *G. lucidum* prolonged the sleeping time of mice by 50% and 60%, respectively, while a 40 mg/kg dose did not significantly increased the sleeping time. In this study, for the first time, the acidic part of the ethanol extract of *G. lucidum* was proven to promote sleeping time by 79.35% (25 mg/kg), 91.03% (50 mg/kg) and 124.37% (100 mg/kg). The sleep promotion effect of GLAA in this study seemed to be better than that of the water extract of *G. lucidum* according to published data, although this conclusion was limited by comparisons between different studies.

The ethanol extract of *G. lucidum* was reported to contain approximately 130–140 triterpene acids, 20 sterols, fewer alkaloids and other compounds. The preparation of GLAA used in our study is also the usual method. GLAA was also found to contain pentobarbital in the present study. Pentobarbital is a well-known sedative-hypnotic and anaesthetic whose hypnotic effect can be enhanced by calcium channel blockers, such as diltiazem and some plant extracts. Calcium ions are widely involved in sleep/wake regulation. Consequently, sleep latency and sleeping time in the pentobarbital-induced sleep test are commonly used as indicators for assessing the sedative and hypnotic effects of dietary supplements.

Many studies have shown that sleep under normal physiological conditions may be influenced by the gut microbiota. PG and LPS are two bacterial cell wall components that are enriched in gram-positive bacteria and gram-negative bacteria, respectively. Our results showed that both PG and LPS levels were altered in serum after gavage administration of GLAA, indicating that GLAA-induced sleep may be related to the gut microbiota. In addition, GLAA may also affect serum LPS/PG levels by regulating gut barrier integrity and preventing translocation into the circulation according to the results of other studies.

GLAA-mediated promotion of sleep in mice. Interestingly, *Bifidobacterium* and *Odoribacter*, which were enriched in the GLAA group, were highly correlated with sleep behaviour and 5-HT in our study. Daidzein can affect various neurobiological regulatory mechanisms such as behaviour, cognition, growth, development and reproduction.Betaine (n-trimethylglycine), a common osmolyte, has function in the modulation of hippocampal neurophysiology and neuroprotection. Interestingly, two herbicides dinitramine and esprocarb were found in these metabolites, although they were not correlated with sleep behaviour or indicators. And we suspected that they might come from the bedding material. Additionally, due to the nontargeted metabolome analysis with LC–MS/MS, integrated database and more included compounds in the database of our study, a large number of metabolites have been explored, which provides us with more ideas about sleep, but these metabolites and results need further verification and research.

Recently, bidirectional interactions between the central nervous system and the gastrointestinal tract have attracted increasing attention. Preclinical evidence supports a role of the gut microbiota in behavioural responses associated with pain, emotion, autism, anxiety, depression, and sleep. The microbiota and the brain communicate with each other via various routes, including the immune system, endocrine and neurocrine pathways, the vagus nerve and the enteric nervous system, involving microbial metabolites such as short-chain fatty acids,
The sleep promotion effects of GLAA were dependent on the gut microbiota. (a) Antibiotics blocked GLAA-induced alterations in sleep behaviour and increases in 5-HT (n = 4–5). (b) Antibiotic treatment did not affect the body weight of mice. (c) Antibiotic treatment decreased microbiota richness, diversity and serum gram-negative bacterial component levels, as evaluated by the Chao1 index, Simpson index and serum LPS content. (d) Composition of the faecal microbiota after antibiotics treatment at the phylum level. The image is generated by R package "ggplot2." (e) Effects of GLAA on the gut microbiota of mice treated with antibiotics. (f) Antibiotics partially depleted gut metabolites altered by GLAA. The data are shown as the means ± SEMs; *p < 0.05, **p < 0.01, ***p < 0.001, ‘ns’ indicates that the difference is not significant. All the data are calculated from 8 to 10 samples per group, and the images are generated by GraphPad Prism 8, except specifically mentioned. GLAA the acidic part of the alcohol extract of G. lucidum mycelia, AB antibiotics, 5-HT 5-hydroxytryptamine, LPS lipopolysaccharide.

Materials and methods

Preparation of GLAA. Ganoderma lucidum CGMCC5.616 was provided by Zhejiang Wuyangtang Pharmaceutical Incorporated Company, Ltd. GLAA was prepared as described with minor modifications. Briefly, 100 g of the G. lucidum fruiting body was crushed, passed through a 20-mesh sieve and mixed with 2 L of 95% ethanol (v/v) before ultrasonic extraction. The solution was ultrasonicated twice for 30 min each time at 40 kHz at room temperature. After filtration, the ethanolic extract was dried by rotary evaporation. The dry product was redissolved in 500 mL of water, degreased with 500 mL of petroleum ether, extracted with 500 mL of ethyl acetate three times, and evaporated to dryness. The ethyl acetate extract was redissolved in 500 mL of ethyl acetate and adjusted to pH 9–10 using 1 mol/L NaOH. Then, the alkalized water layer was isolated and acidified to pH 2–3 with 1 mol/L HCl. The acidified water layer was extracted with ethyl acetate three times to obtain crude GLAA. The GLAA was dried and stored at 4 °C. Experimental research on G. lucidum complied with local and Chinese guidelines and legislation.

Animals. Male specific-pathogen-free (SPF) Institute of Cancer Research (ICR) mice were purchased from Hangzhou Medical College and kept in SPF animal centre of Hangzhou Medical College under controlled light (12 h light–dark cycle), temperature (22–26 °C) and humidity (40–60%) conditions with free access to food and water. Mice that weighed 20–24 g (approximately 6 weeks) were randomly distributed into groups containing ten animals each (five mice per cage) using the standard = RAND() function in Microsoft Excel. The mice were allowed to acclimate to the laboratory conditions for 7 days before pharmacological tests. All experimental procedures were approved by the Animal Experimentation Ethics Committee of Hangzhou Medical College. The experiments were performed in accordance with the criteria of the “Guide for the Care and Use of Laboratory Animals” (NIH publication 86-23, revised 1985) and the ARRIVE guidelines (https://arriveguidelines.org).

Pentobarbital-induced hypnosis test. The pentobarbital hypnosis test was performed as described previously. GLAA was resolved in 0.05% CMC-Na. For the GLAA assay, forty mice were assigned to 4 groups and orally administered GLAA (25, 50 or 100 mg/kg) or 0.05% CMC-Na as a solvent control from 8:00 a.m. to 9:00 a.m. every day for 4 weeks. For the antibiotic depletion experiment, forty mice were assigned to 4 groups: pre-treated saline (control and 100 mg/kg groups) or antibiotics (control + AB and 100 mg/kg + AB groups) for 40 days. On the 13th to the 40th day, the 100 mg/kg + AB group were gavaged with 100 mg/kg GLAA, while the other two groups (control and control + AB) were administered 0.05% CMC-Na.

On the 28th day of the GLAA assay and the 40th day of the antibiotic depletion experiment, 1 h after the last administration of GLAA, 50 mg/kg pentobarbital sodium was injected subcutaneously. The animals were considered asleep if they stayed motionless when positioned on their back and when they lost their righting reflex. The time interval between injection of pentobarbital and start of sleep was considered sleep latency, while prolongation of sleep was described as sleeping time. Faecal samples were collected on the 28th day after administration of GLAA. Mice were sacrificed under carbon dioxide asphyxia 24 h after the pentobarbital-induced hypnosis test. Blood and tissue (brain, spleen, liver, kidney, thymus and colon) were collected. All samples were stored at −80 °C until use.

Antibiotic-induced depletion of the gut microbiota. Antibiotics treatment was performed according to Refs. and divided into two stages. In the first stage, mice were gavaged with amphotericin B (Solarbio Life Science, Beijing, China) at a dose of 1 mg/kg body weight every 12 h for three days. The second stage was from the 4th day to the 40th day, when the water was supplemented with 1 g/L ampicillin (Sangon Biotech, Shanghai, China). Furthermore, an antibiotic combination consisting of 50 mg/kg vancomycin (Sangon Biotech, Shanghai, China), 100 mg/kg neomycin (Sangon Biotech, Shanghai, China), 100 mg/kg metronidazole (Sangon Biotech, Shanghai, China), and 1 mg/kg amphotericin B was administered by gavage every 12 h in the second stage.

Branches-branch chain amino acids, and peptidoglycans. According to our results, 5-HT and the serotonergic synapse pathway might be among the routes connecting the gut microbiota and sleep signals in the brain. In summary, in our study, the effect of the acidic part of the alcohol extract of G. lucidum mycelia on sleep was studied to show that GLAA significantly shortened sleep latency and prolonged sleeping time in pentobarbital-treated mice. This effect was mainly related to an increase in serotonergic synapse signalling in the hypothalamus and was dependent on the gut microbiota of mice.
Figure 5. Gavage administration of GLAA regulated the serotonergic synapse pathway in the hypothalamus. (a) PCA score plots of transcriptome profiles of these four groups. (b) Volcano map of gene transcription after administration of GLAA. (c) Gene transcription was altered after GLAA administration in the four groups ($p_{adj}>0.05$). (d) KEGG pathways altered in the hypothalamus after administration of 100 mg/kg GLAA ($p>0.05$). (e) Gene transcription in the serotonergic synapse pathway was altered after GLAA administration in the four groups. (f) General view of the effects of GLAA on sleep promotion. The data are calculated from 5–6 samples per group. The data are shown as the means ± SEMs; *$p<0.05$, **$p_{adj}<0.05$, 'ns' indicates that the difference is not significant. The images a, b and c are generated by R package "scatterplot3d", "ggplot2" and "pheatmap" respectively, the image f is generated by ScienceSlides Suite, and the others are generated by GraphPad Prism 8. GLAA the acidic part of the alcohol extract of G. lucidum mycelia, AB antibiotics, PCA principal component analysis, 5-HT 5-hydroxytryptamine.
Neurotransmitter and bacterial component detection. Detection of the neurotransmitters 5-hydroxytryptamine (5-HT), γ-aminobutyric acid (GABA), norepinephrine and dopamine (Jianchen, Nanjing, China) in the hypothalamus, as well as detection of LPS (Jianchen, Nanjing, China) and PG (AMEKO, Shanghai, China) in serum, was carried out with ELISA kits according to the manufacturer's instructions.

Gut microbiota analysis. Faecal microbial DNA was extracted using the QIAamp DNA Stool MiniKit (Qiagen, Hilden, Germany). The PCR primers 341F (5′-CCTACGGGNGGCWGCAG-3′) and 805R (5′-GACTACHVGGGTATCTAATCC-3′) targeting the V3–V4 region of the 16S rDNA gene with specific barcodes were used48. PCR was performed in a 25 μL PCR system containing 50 ng of template DNA, 25 pmol of each primer and 12.5 μL of Phusion Hot Start Flex 2X Master Mix (New England Biolabs, Ipswich, MA, USA) with initial denaturation at 98 °C for 30 s, 32 cycles of amplification (denaturation at 98 °C for 10 s; annealing at 54 °C for 30 s and extension at 72 °C for 45 s) and a final extension at 72 °C for 10 min. After purification, sequencing was performed using the Illumina Novaseq platform (Illumina, San Diego, CA, USA).

Raw sequence reads were assigned to each sample according to their unique barcode pairs. Overlapping paired-end reads were merged to form tags using FLASH (version 1.2.11)55. The following quality control criteria were used: (1) an exact match to at least one end of barcodes and primers; (2) no undetermined bases in the tags; and (3) no more than three mismatches in the overlap region. Operational taxonomic units (OTUs) were clustered with a 97% similarity cut-off using Usearch 1156. Alpha diversity was determined using the R packages “rich” and “diversity”. Principal coordinate analysis (PCoA), a standard multivariate statistical technique, was performed with the R package “ade4” to explain differences among microbial communities. The longest sequence in each OTU was chosen as the representative sequence for identification using the Ribosomal Database Project (RDP) database v.11.357. The Chao1 index, Shannon index and principal coordinates analysis (PCoA) of unweighted UniFrac beta diversity were calculated by QIIME software (version 1.8.0) to investigate the richness and diversity of each gut microbiome26. Linear discriminant analysis effect size (LDA) analysis was performed on the Galaxy web platform59. The effect size of each differential taxon was estimated by the linear discriminant analysis (LDA) score, and the taxa with LDA scores greater than 3.5 were defined as discriminative taxa.

Metabolic analysis. LC–MS/MS was used for metabolic analysis of 0.1 g of frozen faeces. A mixture containing 0.1 g of frozen faeces, 300 μL of ice-cold water and 3–4 cold 3 mm steel balls (Jingxin, Shanghai, China) was whirled for 5 min. Then, 500 μL of methanol containing 1 ppm dl-2-chlorophenylalanine was added into the mixture without steel balls, and the mixture was further centrifuged at 12,000 rpm and 4 °C for 10 min. Then, 600 μL of supernatant was sucked into another centrifuge tube and concentrated. Then, 100 μL of 5% methanol water was added to the dried product, mixed and centrifuged at 12,000 rpm at 4 °C for 10 min. Finally, the supernatant was collected for LC–MS/MS (Agilent Technologies Inc., Palo Alto, CA, USA) analysis.

The analytical conditions were as follows: UPLC: column, Waters ACQUITY UPLC HSS T3 C18 (1.8 μm, 2.1 mm × 100 mm); column temperature, 35 °C; flow rate, 0.3 mL/min; injection volume, 1 μL; solvent system, water (0.01% methanolic acid):acetoniitrite; gradient program of positive ion, 95:5 V/V at 0 min, 79:21 V/V at 3.0 min, 50:50 V/V at 5.0 min, 30:70 V/V at 9.0 min, 5:95 V/V at 10.0 min, 95:5 V/V at 14.0 min; gradient program of negative ion, 95:5 V/V at 0 min, 79:21 V/V at 3.0 min, 50:50 V/V at 5.0 min, 30:70 V/V at 9.0 min, 5:95 V/V at 10.0 min, and 95:5 V/V at 14.0 min.

The original data file obtained by LC–MS/MS was first converted to mzML format by ProteoWizard software60. Peak extraction, alignment and retention time correction were performed by the XCMS program61. The “SVR” method was used to correct the peak area. Peaks with deletion rates > 50% were filtered from each group of samples. Then, metabolic identification information was obtained by searching the laboratory’s in-house database and integrating the public database (HMDB and KEGG) and then combining it with metDNA26. Unbiased principal component analysis (PCA), orthogonal partial least squares discriminant analysis (OPLS-DA) and variable importance in projection (VIP) were calculated using SIMCA software (version 14.1) (Sartorius Stedim Biotech, Umeå, Sweden).

Comparative transcriptomic analysis. Total RNA from the hypothalamus was extracted using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) following the manufacturer’s protocol. The total RNA quantity and purity were analysed with a 2100 Bioanalyzer and the RNA 6000 Nano LabChip Kit (Agilent, Palo Alto, CA, USA) with RIN number > 7.0. Poly(A) RNA was purified from total RNA (5 µg) using poly-T oligo-attached magnetic beads using two rounds of purification. Following purification, the mRNA was fragmented into small pieces using divalent cations under elevated temperature. Then, the cleaved RNA fragments were reverse-transcribed into cDNA using oligo(dT) primer (AMERICAN TYPE CULTURE COLLECTION, Rockville, MD, USA) and variable importance in projection (VIP) were calculated using SIMCA software (version 14.1) (Sartorius Stedim Biotech, Umeå, Sweden).

The “SVR” method was used to correct the peak area. Peaks with deletion rates > 50% were filtered from each group of samples. Then, metabolic identification information was obtained by searching the laboratory’s in-house database and integrating the public database (HMDB and KEGG) and then combining it with metDNA26. Unbiased principal component analysis (PCA), orthogonal partial least squares discriminant analysis (OPLS-DA) and variable importance in projection (VIP) were calculated using SIMCA software (version 14.1) (Sartorius Stedim Biotech, Umeå, Sweden).

Linewidth analysis (LDA) score, and the taxa with LDA scores greater than 3.5 were defined as discriminative taxa.

The analytical conditions were as follows: UPLC: column, Waters ACQUITY UPLC HSS T3 C18 (1.8 μm, 2.1 mm × 100 mm); column temperature, 35 °C; flow rate, 0.3 mL/min; injection volume, 1 μL; solvent system, water (0.01% methanolic acid):acetoniitrite; gradient program of positive ion, 95:5 V/V at 0 min, 79:21 V/V at 3.0 min, 50:50 V/V at 5.0 min, 30:70 V/V at 9.0 min, 5:95 V/V at 10.0 min, 95:5 V/V at 14.0 min; gradient program of negative ion, 95:5 V/V at 0 min, 79:21 V/V at 3.0 min, 50:50 V/V at 5.0 min, 30:70 V/V at 9.0 min, 5:95 V/V at 10.0 min, and 95:5 V/V at 14.0 min.

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Comparative transcriptomic analysis. Total RNA from the hypothalamus was extracted using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) following the manufacturer's protocol. The total RNA quantity and purity were analysed with a 2100 Bioanalyzer and the RNA 6000 Nano LabChip Kit (Agilent, Palo Alto, CA, USA) with RIN number > 7.0. Poly(A) RNA was purified from total RNA (5 μg) using poly-T oligo-attached magnetic beads using two rounds of purification. Following purification, the mRNA was fragmented into small pieces using divalent cations under elevated temperature. Then, the cleaved RNA fragments were reverse-transcribed to create the final cDNA library in accordance with the protocol for the mRNA-Seq sample preparation kit (Illumina, San Diego, CA, USA); the average insert size for the paired-end libraries was 300 bp (± 50 bp). Next, we performed paired-end sequencing on an Illumina HiSeq 4000 (Illumina, San Diego, CA, USA) with RIN number > 7.0. Poly(A) RNA was purified from total RNA (5 µg) using poly-T oligo-attached magnetic beads using two rounds of purification. Following purification, the mRNA was fragmented into small pieces using divalent cations under elevated temperature. Then, the cleaved RNA fragments were reverse-transcribed to create the final cDNA library in accordance with the protocol for the mRNA-Seq sample preparation kit (Illumina, San Diego, CA, USA); the average insert size for the paired-end libraries was 300 bp (± 50 bp). Next, we performed paired-end sequencing on an Illumina HiSeq 4000 (Illumina, San Diego, CA, USA) following the vendor's recommended protocol. Then, the sample reads were aligned to the UCSC (http://genome.ucsc.edu/) mouse reference genome using the HISAT package, which initially removes a portion of the reads based on the quality information accompanying each read and then maps the reads to the reference genome. The mapped reads were assembled using StringTie59. Then, all transcriptomes from samples were merged to reconstruct a comprehensive transcriptome using Perl scripts. After the final transcriptome was generated, StringTie and edgeR were used to estimate the expression levels of all transcripts. StringTie was used to determine the expression level of mRNAs by calculating fragments per kilobase of exon model per million mapped reads (FPKM) values64. The differentially expressed mRNAs and genes were selected with log2 (fold change) > 1 or log2 (fold change) < −1 and with statistical significance (P value < 0.05) by the R package.
Statistical analysis. The obtained data were analysed using SPSS 22.0 software (IBM Corporation, Armonk, NY, USA). The data were first tested for normal distribution using the Kolmogorov–Smirnov test. The Mann–Whitney U test (for nonnormal distributions) or Student’s t test (for normal distributions) was performed to test for differences between two groups. Spearman’s rank correlation analysis was conducted using the corplot package of R. The images are generated by GraphPad Prism 8 (GraphPad Software, La Jolla, CA, USA), R and SIMCA software.

Data availability
All data generated or analysed during this study are included in this published article and its additional information files. The data sets generated during the current study are available in the GenBank Sequence Read Archive repository under accession number PRJNA731631.

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Author contributions
Y.Y and L.L. designed this work; C.Y, Z.W, H.X, Y.Z and Y.W performed the experiments; R.Y and H.J. acquired and analysed the data for the work; Y.Y and C.Y drafted the manuscript; L.L. revised the manuscript critically for important intellectual content.

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**Competing interests**
The authors declare no competing interests.

**Additional information**

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**Correspondence** and requests for materials should be addressed to Y.Y. or L.L.

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