

# Specific composition of polyphenolic compounds with fatty acids as an approach in helping to reduce spirochete burden in Lyme disease: *in vivo* and human observational study

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## Abstract

**Background:** Lyme disease (LD) is a tick-borne infection caused by *Borrelia burgdorferi* sensu lato. The current therapeutic approach to this disease is limited to antibiotics. However, after their administration, about 20% of patients experience delayed onset of this illness manifesting as lingering persistent symptoms.

**Methods:** To determine a suitable approach that would help reduce this number, we examined the efficacy of a composition of polyphenolic compounds (baicalein, luteolin, and rosmarinic acid) with fatty acids (monolaurin and cis-2-decenoic acid), and iodine/kelp in a Lyme disease animal model and volunteers.

**Results:** The results showed that 4 weeks of dietary intake of this composition reduced the spirochete burden in animal tissues by about 75%. Basic and differential blood parameters did not show significant differences between control animals and the animals fed with this composition. Also, hepatic and renal toxicity markers were not changed and apoptosis was not observed. Relevant inflammatory cytokines such as IL-6, IL-17, TNF- $\alpha$ , and INF- $\gamma$ , were elevated in infected animals but normalized in infected and treated animals. A small observational study revealed that after administration of this composition to 17 volunteers three times per day for 6 months, 67.4% of the volunteers with late or persistent LD, and not receptive to previous antibiotic application, responded positively, in terms of energy status as well as physical and psychological wellbeing to supplementation with this composition, while 17.7% had slight improvement, and 17.7% were none responsive.

**Conclusion:** We concluded that this specific composition revealed feasible benefits in late or persistent LD management, although double-blind controlled clinical trials are warranted.

**Keywords:** fatty acids, inflammation, Lyme disease, polyphenols, spirochetes

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## Introduction

Lyme disease (LD) is a systemic zoonosis prevalent worldwide.<sup>1,2</sup> It is spread by ticks which transmit bacteria of the genus *Borrelia* while feeding on animals and humans.<sup>3,4</sup> The number of reported LD cases has systematically grown over the past 20 years with the latest estimates reaching 300,000 cases annually in the USA alone.<sup>5</sup> Its causative pathogen, *Borrelia burgdorferi* sensu lato,

is prevalent on the east and west coasts of the USA as well as in the central and eastern parts of Europe. LD affects people of all ages and both genders, although the highest rates have been documented in children aged 10–14 years and in adults over 45 years old.<sup>5–7</sup> The clinical manifestations of LD vary, however common symptoms have been identified. The early signs of LD account for a skin lesion called erythema migrans

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(EM) and/or flu-like symptoms, whereas the systemic symptoms include arthritis, neurologic problems, and cardiac abnormalities which can appear approximately 4–6 weeks after a tick's bite. Persistent fatigue and aches/pain may develop in about 20% of those individuals who followed the recommended antibiotic treatment and can last beyond 6 months. This phenomenon has been described as PTLDS (post-treatment Lyme disease syndrome).<sup>5,8–10</sup>

Several US Food and Drug Administration (FDA)-approved antibiotics are used as primary therapeutics in patients with LD. The first choice for early stages of LD is usually a 2–4-week administration of doxycycline for adults and amoxicillin for children. For late-stage LD, ceftriaxone or cefotaxime are recommended for about the same treatment period. Although some clinical trials have brought contradictory results, it is generally agreed that prolonged antibiotic treatment is not recommended for patients with PTLDS.<sup>5,11,12</sup>

The efficacy of naturally occurring and biologically active substances as anti-borreliae agents is still not well explored, although the number of research investigations with such agents has been growing.<sup>13–16</sup> Our previous *in vitro* study showed that a specific combination of polyphenols with fatty acids and iodine has significant bactericidal effect against two species of *Borrelia* that have been recognized as a pathogenic factor of LD in the USA and Europe. Moreover, this defined composition of phytochemicals worked synergistically and was shown to affect the membrane but not the DNA of the bacteria, demonstrating significant anti-oxidative and anti-inflammatory properties at the same time.<sup>17</sup>

In this study, we report the efficacy of this specific composition of plant-derived compounds against *Borrelia burgdorferi* in an animal model of LD and volunteer patients with a late or persistent form of LD. We attempt here to provide a more comprehensive evaluation of this composition as a potential alternative or perhaps adjunct approach to LD, which needs to be further validated by large double-blind controlled clinical trials.

### Materials and methods

Compounds such as baicalein, luteolin, rosmarinic acid, and cis-2-decenoic acid (10-HAD), with a purity between 90% and 95% according to the manufacturer, were obtained from Baoji GuoKang

Bio-Technology Co. Ltd (Baoji City, China). Organic kelp with standardized iodine content (i.e. 150 µg/ml as 100% of recommended daily allowance, and 60 minerals, vitamins, protein, fats, carbohydrates, and dietary fibers as approximately 25% of daily values) was purchased from Thorvin Inc. (New Castle, VA, USA), and monolaurin as a pure sn-1 monolaurin (glycerol monolaurate) was from Purem Biological (Xi'an, China). Additional purity analysis of compounds to potentially exclude the presence of antibiotics all of which were purchased from Cayman Chemical (Ann Arbor, MI, USA), such as doxycycline and amoxicillin and the anti-inflammatory drug methotrexate used in LD treatment<sup>1,9</sup> was performed at the mass spectroscopy laboratory of Oregon State University and analyzed at the mass spectroscopy laboratory of Stanford University (Supplementary Figures 1Sa–c and Figures 2Sa–d).

### Test microorganisms

A clonal derivative of *Borrelia burgdorferi* B31 strain, MSK5, containing all plasmids, was used in this study.<sup>18</sup> It was cultured in commonly used conditions, that is, Barbour-Stoenner-Kelly H (BSK-H) medium supplemented with 6% rabbit serum (Sigma, St. Louis, MO, USA) without antibiotics at 33°C with 5% CO<sub>2</sub>, in sterile screw-cap 15 ml polystyrene test tubes.

### Test diet

Standard rodent diet was purchased from LabDiet Inc. (St. Louis, MO, USA). The test diet was prepared by LabDiet Inc. and was composed of standard diet enriched with baicalein (650 mg/kg diet), luteolin (300 mg/kg diet), rosmarinic acid (500 mg/kg diet), monolaurin (500 mg/kg diet), 10-HAD (500 mg/kg diet), and iodine 1 mg/kg diet.

### Animal study

A total of 32 C3H/HeN inbred female mice weighing approximately 20 g were obtained from Charles River Laboratories (Wilmington, MA, USA). The 4-week-old mice were housed at the animal facility located at the Dr. Rath Research Institute at an ambient temperature of 21°C. Food and water were provided *ad libitum* during a light and dark cycle of 12 h. Experimental animal protocol No. 01/B052014 was reviewed by and approved in 2014 by the Animal Care and Use Committee

at the Dr. Rath Research Institute. Mice were randomly divided into four experimental groups (eight animals per group): WT (not infected animals fed with the standard diet), LD (infected animals fed with the standard diet), WT + T (not infected animals fed with the test diet), and LD + T (infected animals fed with the test diet). Low passage of virulent *Borrelia burgdorferi*, grown at 33°C in a BSK-H medium until reaching a concentration of approximately  $1 \times 10^7$  spirochetes/ml, were inoculated intradermally ( $10^4$  spirochetes/mouse) into animals from two experimental groups, LD and LD + T, as previously reported.<sup>19</sup> Mice from groups WT and WT + T received mock injections of  $1 \times$  phosphate buffered saline (PBS). After 3 weeks, mice from the WT + T and LD + T groups were transferred from the standard diet to the test diet and fed for a subsequent 4 weeks. The rest of the animals were kept on the standard diet. At the end of the study all animals were sacrificed and the tissue samples, including blood, were collected for further testing. Control animals were mice fed with the standard diet and not infected (WT) as well as mice fed with the standard diet and infected with *Borrelia burgdorferi* (LD). Blood was aseptically obtained from all experimental mice by an intracardiac puncture using a 1-ml tuberculin syringe with a 27-G needle and separated into two samples. One sample of the collected blood was used for basic and differential hematology, and the second sample was used to prepare serum for biochemical analysis. Tissue samples from the ears, heart base, tibiotarsal joints, liver, kidneys, and spleen were collected, immediately snap-frozen in liquid nitrogen, and subjected to quantitative polymerase chain reaction (qPCR Bio-Rad, Hercules, CA, USA) analysis and/or immunohistochemistry. In addition, blood and tissue cultures (from the ear, heart base, spleen, and joints) were processed and scored for spirochetal growth in BSK-H medium after 3–5 weeks as previously described.<sup>19</sup>

#### Quantitative analysis of *Borrelia burgdorferi* DNA

To confirm the presence of infection and quantify spirochete tissue burdens, all samples were analyzed by qPCR as previously reported.<sup>18,19</sup> Briefly, whole genomic DNA was extracted from tissue samples (ears, heart base, spleen, and tibiotarsal joints) using DNeasy Blood and Tissue kit (Qiagen, Valencia, CA, USA) according to the manufacturer's instructions. Before DNA extraction, all

tissue samples were snap-frozen in liquid nitrogen, weighed, and homogenized using a QIA shredder kit (Qiagen). Optimized primer sets of *Borrelia burgdorferi* flagellin (*flaB*, *bb0147*) gene (F-5' TCTTTTCTCTGGTGAGGGAGCT; R-5' TCC TTCCTGTTGAACACCCTCT) and murine  $\beta$ -actin (*ACTB*) (F-5' CAAGTCATCACTATTGG CAACGA; R-5' CCAAGAAGGAAGGCTGGAA AA) gene from published sequences were used.<sup>19</sup> All qPCR reagents were purchased from Qiagen. DNA templates [i.e.  $1 \times 10^5/\mu\text{l}$  *B. burgdorferi* equivalents from American Type Culture Collection (ATCC) (Manassas, VA, USA) and mouse from Promega (Madison, WI, USA) served as known amounts of spirochetal and mouse genomic DNA], were included in each qPCR experiment, and used to compile the standard curves, and to determine the total number of spirochetes (*flaB* copy number) in collected mouse tissue samples. Also, a no-template reaction control was included. QuantiTect SYBR Green PCR Kit (Qiagen) was utilized to determine the borrelial *flaB* target gene, expressed as per 1 mg of tissue weight. It contained QuantiTect SYBR Green PCR Buffer, pH=8.7, as the source of 2.5 mM  $\text{Mg}^{2+}$ , deoxynucleotides and HotStarTaq DNA polymerase, in addition to each primer used at a concentration of 0.3  $\mu\text{M}$  in a 50  $\mu\text{l}$  qPCR mixture. The qPCR cycler conditions (BioRad CFX96) were programmed according to manufacturer's recommendations and were as follows: an activation step at 95°C for 15 min, 40 cycles of: denaturation at 94°C for 15 s, annealing at 59°C for 30 s, and extension at 72°C for 30 s. To verify the specificity and identity of the qPCR product, melting curve analyses were performed. Additional validation was conducted by subjecting the tissue samples to an independent qPCR analysis performed by a certified laboratory (IDEXX BioResearch, Sacramento, CA, USA). Each sample was analyzed in triplicate.

#### Blood hematology

Blood samples obtained from all experimental mice were submitted to a certified analytical laboratory (IDEXX BioResearch) where the basic (RBC, Red Blood Cells; WBC, White Blood Cells; HGB, Hemoglobin; HCT, Hematocrit; MCV, Mean Corpuscular Volume; MCH, Mean Corpuscular Hemoglobin; and MCHC, Mean Corpuscular Hemoglobin Concentration) and differential (neutrophils, monocytes, lymphocytes, and platelets counts/levels) hematologic analysis was performed and supported by

pathological evaluation. Each sample was analyzed in triplicate.

#### *Biochemical analysis*

Sera from all experimental animals were analyzed by a certified analytical laboratory (IDEXX BioResearch) for alkaline phosphatase (ALP), alanine aminotransferase (ALT), aspartate aminotransferase (AST), gamma-glutamyl transferase (GGT), creatinine, cholesterol, glucose, blood urea nitrogen (BUN), bilirubin, haptoglobin, and kidney injury molecule-1 (KIM-1) levels. KIM-1, a 30 KDa, type 1 membrane protein, has been considered as a potential biomarker for renal injury since it is highly upregulated in the proximal tubule of the injured kidney, but exists in very low levels in normal uninjured kidneys.<sup>20</sup> In addition, 50  $\mu$ l serum from each mouse was subjected to pro-inflammatory cytokines and chemokines analysis using the Analyte Profiler ELISArray assay kit (SA Biosciences, Germantown, MD, USA) according to the manufacturer's protocol. Each sample was analyzed in triplicate.

#### *Immunohistochemistry*

Samples from kidney, liver, and joints, collected at the end of the experiment, were fixed in 10% neutral-buffered formalin for 24–48 h and subjected to immunohistochemistry evaluation performed by an independent certified laboratory (HistoTox Labs, Boulder, CO, USA). The 5  $\mu$ m sections, placed on glass slides, were stained with hematoxylin and eosin, anti-F4/80 (macrophage marker), and anti-ACS3 (activated caspase-3, apoptotic marker) antibodies. Caspase-3 is activated by the upstream caspase-8 and caspase-9, serving as a convergence point for intrinsic and extrinsic apoptotic signaling pathways. Thus activation of the caspase-3 pathway is a hallmark of apoptosis and can be used in cellular assays to quantify activators and inhibitors of the 'death cascade'.<sup>21</sup>

F4/80, also known as EGF-like module-containing mucin-like hormone receptor-like 1 (Epidermal Growth Factor-like module-containing mucin-like hormone receptor-like 1) is a member of the adhesion G-protein coupled receptor (GPCR) family. F4/80 antigen is a mature mouse cell surface glycoprotein expressed at high levels on various macrophages including: Kupffer cells, splenic red pulp macrophages, microglia, gut lamina propria, and Langerhans cells in the skin. F4/80

antigen is also expressed on the macrophages of the connective tissue, heart, kidney, and reproductive and neuroendocrine systems, thus it is a well-known and widely used marker of murine macrophage populations.<sup>22</sup> Sections were cryptically coded and an additional unbiased histopathological examination was performed by an onsite board-certified pathologist.

#### *Human observational study*

Between 2014 and 2015, eligible adult patients ( $n=17$ ) were recruited through referrals from physicians in Germany where the study was conducted at the Private Praxisklinik H. Baltin, Bavaria, Germany. The study followed the protocol adhering to the regulations of the 1975 Declaration of Helsinki and was approved by the institutional Ethics Committee (No. DRRI/2014) at the participating center (Private Praxisklinik H. Baltin, Bavaria, Germany), and supported by written informed consent from each patient. Patients were classified as eligible for the study if they were at least 18 years old, with a history of acute LD acquired in Germany, and with at least one of the following: a history of single or multiple EM, neurologic symptoms attributed to LD, or Lyme arthritis. Additional medical documentation proving that the patients currently suffered from late or chronic LD (manifesting in the form of fatigue, musculoskeletal pain, arthritis, cardiac arrhythmia, neuropathies, cognitive dysfunction/paralysis) and that they previously had undergone treatments with a recommended antibiotic regimen was mandatory.<sup>5,23</sup> At the time of enrollment, all patients had the following symptoms that interfered with their regular functioning and ability to work: widespread musculoskeletal pain and/or cognitive impairment with accompanying profound fatigue. The chronic symptoms had to have begun within 6 months after the initial application of the recommended antibiotic regimen and had to have persisted beyond 6 months.<sup>23</sup> Patients received the test composition in the form of capsules containing baicalein 250 mg/day, luteolin 75 mg/day, rosmarinic acid 100 mg/day, monolaurin 250 mg/day, 10-HAD 100 mg/day, and iodine 0.15 mg/day (in the form of kelp), three times per day for 6 months. During the oral treatment phase, compliance and safety were monitored through monthly patient visits at the doctor's clinic. Clinical and laboratory evaluations were performed on day 90 and day 180 and included a complete medical history, a detailed physical examination,

**Table 1.** Summary of mice physiological parameters after 4 weeks of treatment.

Parameters	WT (n=8)	LD (n=8)	WT + T (n=8)	LD + T (n=8)
RBC ( $\times 10^6/\text{mm}^3 \pm \text{SD}$ )	8.11 $\pm$ 2.2	7.48 $\pm$ 2.6	8.56 $\pm$ 2.8	8.78 $\pm$ 2.5
WBC ( $\times 10^3/\text{mm}^3 \pm \text{SD}$ )	6.3 $\pm$ 1.9	6.8 $\pm$ 2.3	6.1 $\pm$ 2.4	6.4 $\pm$ 2.2
HGB (g/L $\pm$ SD)	142 $\pm$ 23	146 $\pm$ 27	147 $\pm$ 35	134 $\pm$ 39
HTC (% $\pm$ SD)	41.4 $\pm$ 6.5	43.2 $\pm$ 5.6	41.8 $\pm$ 5.8	41.9 $\pm$ 7.6
MCV (fL $\pm$ SD)	49.3 $\pm$ 7.5	49.3 $\pm$ 5.9	46.0 $\pm$ 6.7	49.4 $\pm$ 6.9
MCH (pg $\pm$ SD)	14.8 $\pm$ 5.9	13.4 $\pm$ 7.1	14.9 $\pm$ 5.8	14.6 $\pm$ 5.6
MCHC (mmol/L $\pm$ SD)	18.9 $\pm$ 4.2	18.9 $\pm$ 4.2	20.6 $\pm$ 4.3	20.0 $\pm$ 4.7
Neutrophils (% $\pm$ SD)	25.8 $\pm$ 6.8	29.6 $\pm$ 7.5	24.4 $\pm$ 6.7	24.5 $\pm$ 7.9
Monocytes (% $\pm$ SD)	1.9 $\pm$ 0.6	4.1 $\pm$ 0.5*	1.6 $\pm$ 0.2	2.7 $\pm$ 0.3
Lymphocytes (% $\pm$ SD)	66.3 $\pm$ 8.6	84.9 $\pm$ 7.5	74.7 $\pm$ 9.1	75.8 $\pm$ 8.2
Platelets	Adequate	Adequate	Adequate	Adequate

\* $p < 0.01$ ; HGB, hemoglobin; HTC, hematocrit; LD, mice infected with *Borrelia burgdorferi* (intradermal injection with  $10^4$  spirochetes/mouse); LD + T, mice infected with *Borrelia burgdorferi* (intradermal injection with  $10^4$  spirochetes/mouse) and fed with plant-based diet for 4 weeks; MCH, mean corpuscular hemoglobin; MCHC, mean corpuscular hemoglobin concentration; MCV, mean corpuscular volume, RBC, red blood cells; SD, standard deviation; WBC, white blood cells; WT; control mice; WT + T, control mice fed with plant-based diet for 4 weeks.

neuropsychological testing, and sampling of peripheral blood, if deemed necessary.

### Statistical analysis

All the data are presented as means  $\pm$  standard deviation (SD) ( $n=8$  for the animal study, and  $n=17$  for the human study). Student's two-tailed  $t$  test was used to determine statistically significant differences set at 0.05 levels. Statistical analysis was performed using GraphPad software.

## Results

### Effect of the test composition in vivo

After completion of 4 weeks of treatment with the test diet, there were no significant variations in the weight as well as food and water consumption between all experimental groups (Table 1). Evaluation of blood revealed no alterations in hematology parameters between experimental groups, except that there was a significantly increased level of monocytes in the animals infected with *Borrelia burgdorferi* (LD) (Table 2). Also in the

same animal group an increased level of neutrophils was noticed, but it did not reach statistical significance ( $p=0.053$ ). Pathologist analysis further revealed no signs of hemolysis and lipemia, nor any anisocytosis or poikilocytosis. Also, the level of (meta/pro)-myelocytes was classified as 'undetectable'. Furthermore, biochemical analysis of sera from all experimental animals showed no changes in the levels of liver enzymes and basic clinical metabolic markers such as cholesterol, glucose, bilirubin, BUN, and creatinine. The LD group displayed an increased level of ALP, however, it was not statistically significant ( $p=0.055$ ) (Table 3). In addition, haptoglobin and KIM-1 levels showed no changes, and pathologist evaluation reported no toxicity in the tissues (Figure 1).

Pro-inflammatory cytokines such as IL-6, IL-17, INT- $\gamma$ , and TNF- $\alpha$ , were elevated in the LD group, however their levels were normalized in the LD + T group (Figure 2). None of the control groups (WT and WT + T) tested for *Borrelia burgdorferi* DNA showed detectable levels. In the LD group of mice, the borrelial DNA was detected in the ears, heart base, spleen, and joints

**Table 2.** Basic and differential blood parameters of mice after 4 weeks of treatment.

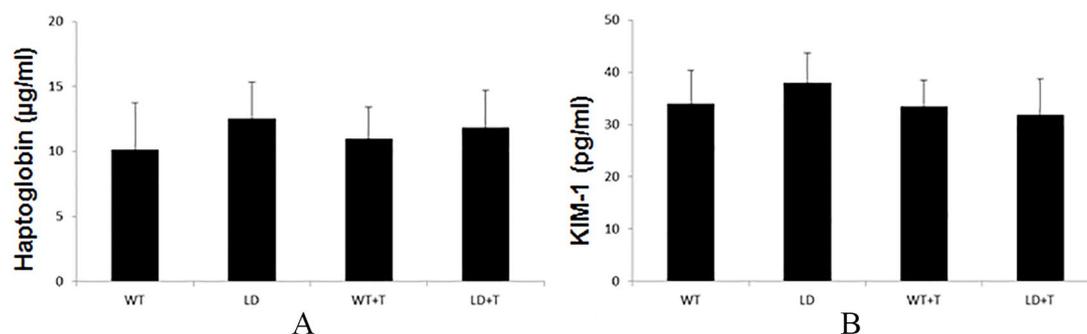
Parameters	WT (n=8)	LD (n=8)	WT + T (n=8)	LD + T (n=8)
Weight (g/mouse $\pm$ SD)	25.3 $\pm$ 3.1	24.3 $\pm$ 2.9	25.2 $\pm$ 3.1	25.1 $\pm$ 3.1
Food (g/mouse/day $\pm$ SD)	3.3 $\pm$ 0.6	3.2 $\pm$ 1.1	3.1 $\pm$ 1.0	3.4 $\pm$ 0.8
Water (ml/mouse/day $\pm$ SD)	2.4 $\pm$ 0.3	2.6 $\pm$ 0.7	2.8 $\pm$ 0.8	2.5 $\pm$ 0.4

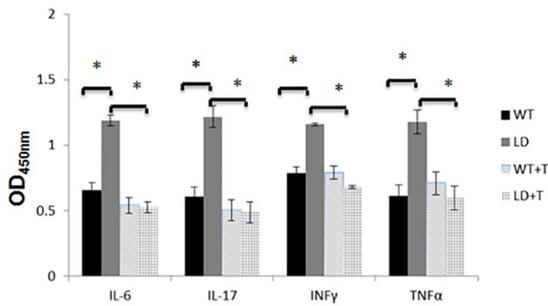
LD, mice infected with *Borrelia burgdorferi* (intradermal injection with 10<sup>4</sup> spirochetes/mouse); LD + T, mice infected with *Borrelia burgdorferi* (intradermal injection with 10<sup>4</sup> spirochetes/mouse) and fed with plant-based diet for 4 weeks; SD, standard deviation; WT, control mice; WT + T, control mice fed with plant-based diet for 4 weeks.

**Table 3.** Biochemical parameters of mice sera after 4 weeks of treatment.

Parameters	WT (n=8)	LD (n=8)	WT + T (n=8)	LD + T (n=8)
ALP (U/L $\pm$ SD)	83.1 $\pm$ 8.7	100.9 $\pm$ 7.6	89.1 $\pm$ 7.3	70.9 $\pm$ 7.9
ALT (U/L $\pm$ SD)	40.3 $\pm$ 5.5	35.7 $\pm$ 4.9	31.3 $\pm$ 5.9	36.9 $\pm$ 7.6
AST (U/L $\pm$ SD)	182.5 $\pm$ 10.2	257.8 $\pm$ 14.3	162.7 $\pm$ 11.5	141.9 $\pm$ 9.9
GGT (U/L)	<3	<3	<3	<3
Total bilirubin ( $\mu$ mol/L)	<3.4	<3.4	<3.4	<3.4
Cholesterol (mmol/L $\pm$ SD)	3.8 $\pm$ 0.3	3.8 $\pm$ 0.3	3.9 $\pm$ 0.3	3.3 $\pm$ 0.2
Glucose (mmol/L $\pm$ SD)	6.3 $\pm$ 0.6	6.5 $\pm$ 0.8	5.9 $\pm$ 0.7	5.8 $\pm$ 0.8
BUN (mmol/L $\pm$ SD)	8.6 $\pm$ 1.1	8.2 $\pm$ 0.8	7.9 $\pm$ 0.9	9.5 $\pm$ 1.1
Creatinine ( $\mu$ mol/L)	<26.5	<26.5	<26.5	<26.5

ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; BUN, blood urea nitrogen; GGT, gamma-glutamyl transferase; LD, mice infected with *Borrelia burgdorferi* (intradermal injection with 10<sup>4</sup> spirochetes/mouse); LD + T, mice infected with *Borrelia burgdorferi* (intradermal injection with 10<sup>4</sup> spirochetes/mouse) and fed with plant-based diet for 4 weeks; SD, standard deviation; WT, control mice; WT + T, control mice fed with plant-based diet for 4 weeks.

**Figure 1.** Levels of haptoglobin (a) and KIM-1 (b) in mice sera after completion of 4 weeks of treatment. n=8. KIM-1, kidney injury molecule-1; LD, mice infected with *Borrelia burgdorferi* (intradermal injection with 10<sup>4</sup> spirochetes/mouse); LD + T, mice infected with *Borrelia burgdorferi* (intradermal injection with 10<sup>4</sup> spirochetes/mouse) and fed with plant-based diet for 4 weeks; WT, control mice; WT + T, control mice fed with plant-based diet for 4 weeks.



**Figure 2.** OD readings of mice serum cytokines after completion of the 4-week treatment.  $n=8$ ,  $*p < 0.001$ . IL, interleukin; INT- $\gamma$ , interferon gamma; LD, mice intradermally infected with  $10^4$  spirochetes/mouse and not fed with plant-based diet; LD + T, mice infected intradermally with  $10^4$  spirochetes/mouse and fed for 4 weeks with plant-based diet; OD, optical density; TNF- $\alpha$ , tumor necrosis factor alpha; WT, mice not infected and not fed with plant-based diet; WT + T, mice not infected and fed for 4 weeks with plant-based diet.

of all mice. In the LD + T group borrelial DNA was found only in the joints of 2/8 animals (25%) and at a significantly decreased level compared with the LD group (Table 4 and Figure 3). Results were confirmed by an independent qPCR examination performed by a certified outside laboratory. Immunohistochemistry analysis supported by the pathological evaluation revealed no signs of apoptosis in the liver and kidneys of animals in any of the experimental groups (Figure 4). However, a noticeable increase in staining with anti-F4/80 antibodies was detected in the joints of mice in the LD group, as well as in two animals from the LD + T group in which borrelial DNA was detected, although at markedly reduced intensity (Figure 5).

#### Effect of the test composition in human subject

Evaluation of the dietary supplementation with the six plant-based agents in 17 patients affected by LD revealed that three (3/17) patients became completely symptom-free after 6 months. The patients reported physical and psychological improvements of their symptoms and were able to resume working. Eight patients (8/17) also experienced significant physical and psychological improvement and were able to work again; however, further evaluation was scheduled to follow up on whether the improvements would change after discontinuation of the treatment. Two patients (2/17) experienced only slight improvements, while four patients (4/17) showed no

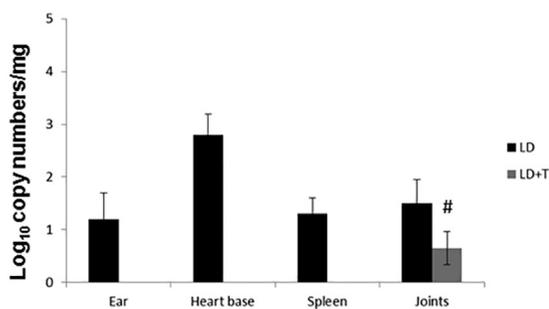
**Table 4.** Comparison of culture and *flaB* DNA qPCR for individual infected mice tissue samples collected after 4 weeks of treatment.

LD culture*	Ear	Heart base	Spleen	Joints
	LD qPCR			
+	+	+	+	+
+	+	+	+	+
+	+	+	+	+
+	+	+	+	+
+	+	+	+	+
+	+	+	+	+
+	+	+	+	+
+	+	+	+	+
LD + T culture*	LD + T qPCR			
-	-	-	-	-
-	-	-	-	+
-	-	-	-	-
-	-	-	-	-
-	-	-	-	-
-	-	-	-	+
-	-	-	-	-
-	-	-	-	-

qPCR, quantitative polymerase chain reaction; LD, mice infected with *Borrelia burgdorferi* (intradermal injection with  $10^4$  spirochetes/mouse); LD + T, mice infected with *Borrelia burgdorferi* (intradermal injection with  $10^4$  spirochetes/mouse) and fed with plant-based diet for 4 weeks,  $n=8$ . \*Culture of ear, heart base, spleen, and joints tissue. (-), *flaB* DNA-negative result in all subjected tissues; (+), *flaB* DNA-positive result in all subjected tissues.

improvement and reported abnormal tiredness, pain in the limbs, headache, and visual disturbances. After discontinuing the supplementation for 2 weeks, these symptoms subsided. One of these four patients (1/4) made a new attempt and resumed the supplementation starting with a lower dose of one capsule per day gradually

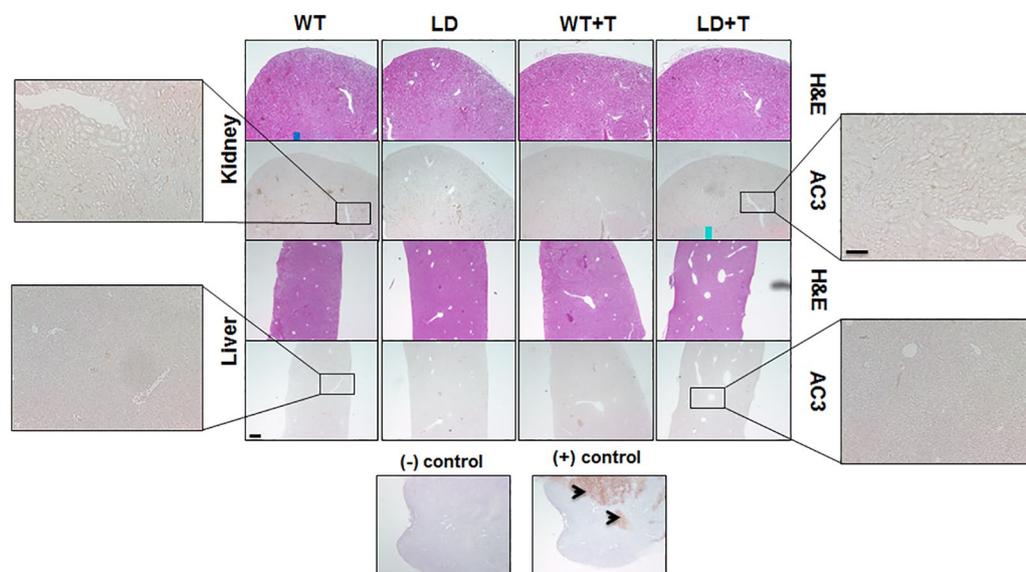
increasing to three capsules per day, which resulted in no negative effects observed with a slight improvement in physical condition. Further diagnostics for co-infections often associated with LD as well as neurological issues with a change in therapy approach were necessary for patients with no improvement, and that took place outside of this observation study (Table 5).



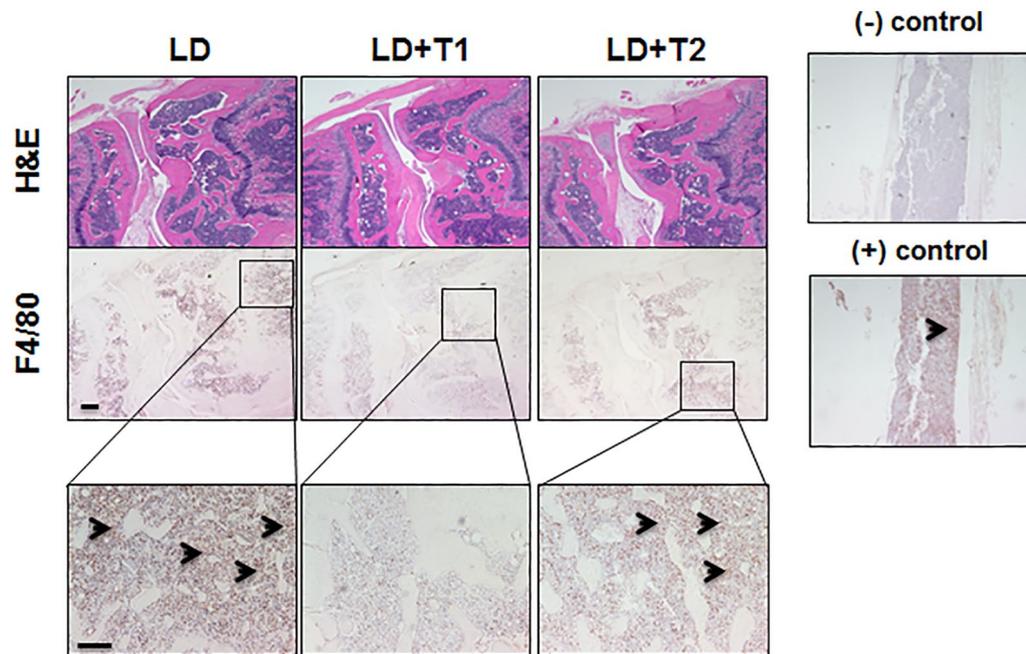
**Figure 3.** Estimation of spirochete burden by quantitative polymerase chain reaction in mice per mg of tissue after completion of 4 weeks of treatment.  $n=8$ ,  $^{\#}p < 0.05$  compared with LD group.  $C_T$  values  $< 40$  were considered positive. LD, mice infected intradermally with  $10^4$  spirochetes/mouse and not fed with plant-based diet; LD + T, mice infected intradermally with  $10^4$  spirochetes/mouse and fed for 4 weeks with plant-based diet.

### Discussion

Here we demonstrated the results from animal and human observational studies after the application of the test composition consisting of six compounds such as baicalein, luteolin, rosmarinic acid, monolaurin, 10-HAD, and kelp/iodine. The present research extends the findings from our previous study demonstrating significant *in vitro* efficacy of this composition against pleomorphic forms of *Borrelia burgdorferi* s.s. and *B. garinii*.<sup>17</sup> We found there that the effect of this combination was comparable to *in vitro* effectiveness of the triple combination of antibiotics (i.e. doxycycline with daptomycin and cefoperazone), and it demonstrated anti-oxidative and anti-inflammatory effects at the same time.<sup>17,24</sup> Results from the animal study concurred with observations on human subjects demonstrating the general safety of this composition when applied in a diet. They revealed no significant changes in basic and differential hematology. Only the infected, but not treated mice (LD group), had an increased level of monocytes and borderline elevated neutrophils, however, these parameters were normalized in animals infected and fed with the test composition (LD + T). Also clinical physiological and biochemical parameters in sera of the experimental groups of animals showed no significant variability.



**Figure 4.** Representative images of AC3 detection in mouse kidney and liver tissues by immunohistochemistry using rabbit polyclonal antibody. Magnifications 4 $\times$ , 10 $\times$  for inserts; [-] negative and [+] positive control of mouse tumor. AC3, active caspase-3; H&E, hematoxylin and eosin staining; LD, mice infected intradermally with  $10^4$  spirochetes/mouse and not fed with plant-based diet; LD + T, mice infected intradermally with  $10^4$  spirochetes/mouse and fed for 4 weeks with plant-based diet; WT, mice not infected and not fed with plant-based diet; WT + T, mice not infected and fed for 4 weeks with plant-based diet.



**Figure 5.** Representative images of anti-F4/80 antibody detection in mice joints tissues by immunohistochemistry using rabbit polyclonal antibody. Magnifications 4×, 10× for inserts; (-) negative and (+) positive control of mouse bone. H&E, hematoxylin and eosin staining; LD, mice intradermally with  $10^4$  spirochetes/mouse and not fed with plant-based diet; LD + T1, mouse no. 2 infected intradermally with  $10^4$  spirochetes/mouse and fed for 4 weeks with plant-based diet with increased staining with anti-F4/80; LD + T2, mouse no. 1 infected intradermally with  $10^4$  spirochetes/mouse and fed for 4 weeks with plant-based diet with no increased staining with anti-F4/80.

**Table 5.** Summary of the observational study of 17 patients with late or chronic LD after 6 months of supplementation with the test composition.

No. patient	MS	SS	PSY/US	No improvement	Slight improvement	Marked improvement	Complete improvement
1	+++	+++	+	-	x	-	-
2	+++	+++	-	x	-	-	-
3	-	+	+++	-	-	-	x
4	+++	+	+++	-	x	-	-
5	++	++	++	-	-	x	-
6	+++	+++	++	x	-	-	-
7	+++	+++	++	x	-	-	-
8	+	+	+++	-	-	x	-
9	+++	++	+	x	-	-	-
10	+++	+++	++	-	-	-	x

(Continued)

**Table 5.** (Continued)

No. patient	MS	SS	PSY/US	No improvement	Slight improvement	Marked improvement	Complete improvement
11	+++	++	-	-	-	-	x
12	+++	+	+++	-	-	x	-
13	+++	++	+	-	-	x	-
14	+++	+++	++	-	-	x	-
15	+++	+	+	-	-	x	-
16	+++	-	+	-	-	x	-
17	+++	++	+++	-	-	x	-

Score: +++, severe symptoms, unable to work; ++, less severe symptoms, ability to work not permanently restricted; +, intermittent slight symptoms.  
LD, Lyme disease; MS, main symptoms: lack of energy/tiredness/low stamina/fatigue/ musculoskeletal pain/forgetfulness/ concentration disorder; PSY/US, psychological symptoms/unusual symptoms: depression/irritability/panic and anxiety attacks/aggression/neuropathies/paralysis; SS, subsidiary symptoms: vertigo/visual disorders/cardiac arrhythmia.

There was an observed increase in inflammatory cytokine levels such as IL-6, IL-17, INT- $\gamma$ , and TNF- $\alpha$ , in the LD group which corroborates other reports and points to ongoing chronic inflammation and might signal ongoing pathophysiological processes.<sup>25</sup> However, the cytokine levels were normalized in the LD + T group of mice which consumed the test diet. Consumption of the diet enriched with the test composition also did not cause toxic effects in mice tissues since the hepatic and renal injury markers were not elevated in the control (WT + T) and the infected (LD + T) animals. This was confirmed by activated caspase-3 and pathologist evaluation of the results.

The infected animals (LD group) showed the presence of detectable borrelial DNA in different tissues, which is consistent with previously published data. However, it is worth mentioning that an *in vivo* study published by Pahl *et al.* and others showed that dissemination of *Borrelia burgdorferi* does not proceed evenly and its distribution can vary among tissues of infected animals.<sup>26-29</sup> Our qPCR results indicated that very low copy numbers of spirochetal DNA were present in the joint tissues from two infected mice fed with the test diet; however, they could not be cultured and were significantly decreased compared with the LD group. The issue of detection of noncultivable *Borrelia burgdorferi* after antibiotic treatment *in vitro* and *in vivo* is not new and was reported previously.<sup>30,31</sup> Interestingly, an increased level of F4/80

marker in the joint tissue was found in the same two mice, although, likewise, at a visibly reduced level compared with the LD group. These results cannot exclude the possibility that spirochetes could retain a low level of infectivity and be a cause of inflammation, but their ability to replicate was altered. Potentially, the retained ability to express borrelial lipoproteins could contribute, since it has been shown that they can potentiate a plethora of pro-inflammatory responses.<sup>32-36</sup> Therefore, the fate of *Borrelia burgdorferi* and its infectivity aspects in the joints needs to be established. However, the presence of enhanced inflammatory areas in the analyzed tissue sections could correspond to the presence of spirochetes in the persistent stage, which should be confirmed by more detailed study. Joint tissue is extracellular matrix (ECM)-rich and, similar to bone, it is difficult to penetrate by antibacterial agents. The reports about the existence of live spirochetes after antibiotic treatment implicating collagen as a key factor of persistence contributing to antibiotic treatment failure have been reported earlier.<sup>37-40</sup>

Treatment with different FDA-approved antibiotics on small and large mammals also seems to reduce the *Borrelia burgdorferi* burden but fails to clear the infection as also documented by many other research groups.<sup>41-50</sup> These *in vivo* reports were further substantiated by clinical trials findings.<sup>5,11,12,46,49</sup> It is also worth noting that a declining antibody titer has been observed after antibiotic

treatment, despite the presence of low levels of persisting spirochetes.<sup>41,50</sup> However, others reported that after antibiotic therapy in dogs, the antibody titers in some animals remained at constant levels, which would argue for the persistence of the antigenic stimulus rather than complete elimination of pathogens.<sup>51</sup>

It is being increasingly accepted that conventional therapy with antibiotics for late and persistent LD brings limited results. Monotherapy especially has minimal long-term success and, although it is still used there has been no solid evidence documenting its efficacy in late and persistent stages of LD to date.<sup>5,11,12</sup> Our previously published *in vitro* results showed that a composition of baicalein, luteolin, rosmarinic acid, monolaurin, 10-HAD, and iodine at their 1/8 minimum inhibitory concentration (MIC) values has significant synergistic effect against *Borrelia* spp. This composition revealed anti-oxidative properties affecting the membrane of *Borrelia* but not its DNA. We also noticed its inhibitory effect on the release of IL-1 $\alpha$ , IL-1 $\beta$ , and IL-6 by human CD14<sup>+</sup> monocytes stimulated with live *Borrelia* sp.<sup>17</sup> To validate these results we performed a small observational study limited, however, to number of patients and clinical and laboratory evaluations. In our pilot study of 17 patients with LD who had previously undergone several rounds of antibiotic treatment but whose health had not improved after the finished therapy, a 6-month intake of a specific combination of phenolic compounds with fatty acids and kelp/iodine improved the health conditions of 64.7% of the patients allowing them to conduct their normal daily activities and resume working. Nonetheless, observed results, although encouraging, need to be further confirmed in a larger double-blind controlled clinical study to ensure that patients improved health conditions are also significant and consistent in a larger population of patients with late or chronic LD, and substantiated by more in-depth clinical evaluation. Only then, may a more confident conclusion about the efficacy of this composition in humans be drawn. Perhaps further health benefits may be observed with an increased dosage and longer time of intake of the composition. However, as the study indicated the dose adjustment should be made gradually. This study also revealed that those patients with little or no improvement should be subjected to further diagnosis for accompanying infections and neurologic dysfunction to exclude the lack of response as the reason.

In summary, this study documented the efficacy of the oral intake of a composition composed of six biologically active plant-derived agents in an animal model of LD and volunteer patients with late or chronic LD. Further studies investigating their efficacy as an adjunct to antibiotic treatment, and as well as in a larger controlled clinical setting, seems to be reasonable and merits further study. Also, it should be mentioned that the natural compounds used in our *in vivo* study reached 90–95% purity, although additional testing performed by us showed similar or even greater purity and lack of antibiotics and anti-inflammatory agents relevant in LD treatment. However, the presence of small amounts of residues could be influential and in a limited way affect experimental outcomes.

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### Author contributions

AG conceived, designed, and performed the *in vivo* experiments, analyzed data, wrote the paper and had primary responsibility for final content; GG and HB designed and performed the human study, validated and had primary responsibility for final content; AN and MR validated, wrote the paper, and had primary responsibility for final content. All authors read and approved the final manuscript.

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### Supplemental material

Supplemental material for this article is available online.

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