

IgM-enriched solution BT086 improves host defense capacity and energy store preservation in a rabbit model of endotoxemia

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Conflicts of interest

The authors state that they have no conflict of interest to declare.

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Introduction: The therapeutic value of intravenous immunoglobulin (IVIG) as an adjuvant therapy in sepsis remains debatable. We hypothesized that intravenous administration of BT086, a predominantly IgM IVIG solution, would improve host defense in an established rabbit model of endotoxemia and systemic sepsis.

Methods: New Zealand white rabbits were randomized into the following four groups: (1) the negative control group without lipopolysaccharide (LPS, control), (2) the positive control group with LPS infusion (LPS group), (3) the albumin-treated LPS group (ALB+LPS group), and (4) the BT086-treated LPS group (BT086 + LPS group). A standardized amount of *E. coli* was intravenously injected into all of the animals. The vital parameters, the concentration of *E. coli* in the blood and other organs, the residual granulocyte phagocytosis activity, and the levels of the inflammatory mediators were measured. Histological changes in the lung and liver tissue were examined following autopsy.

Results: The elimination of *E. coli* from the bloodstream was expedited in the BT086-treated group compared with the LPS- and albumin-treated groups. The BT086 + LPS group exhibited higher phagocytic activity of polymorphonuclear neutrophils (PMNs) than the control and ALB+LPS groups. The liver energy stores were higher in the BT086 + LPS group than in the other groups.

Conclusion: Our data suggest that the IgM-enriched IVIG has the potential to improve host defense in a rabbit model of endotoxemia. Studies using different animal models and dosages are necessary to further explore the potential benefits of IgM-enriched IVIG solutions.

Editorial comment: what this article tells us

The use of IgM-enriched Intravenous Immunoglobulin (IVIG) in the treatment of sepsis remains controversial. In this rabbit model of LPS-induced endotoxemia, administration of an IgM-enriched IVIG solution improved host defense against *E. coli* bacteremia. The findings suggest that the IgM-enriched IVIG solution may improve host defense by neutralizing LPS.

Despite advances in intensive care medicine, the number of sepsis cases continues to increase, and mortality associated with sepsis remains high.¹ Bacterial surface proteins and toxins are considered to be key activators of the complex inflammatory cascade in sepsis.² Antibiotic treatment does not influence the existing bacterial endotoxin load in an animal model of sepsis³ and in humans⁴ and pathogen disintegration followed antibiotics might even aggravate endotoxin release from bacteria.^{3,4} Intravenous immunoglobulin (IVIG) has the potential to scavenge and neutralize endotoxins, which reduces the overall pro-inflammatory reaction.⁵ The IgM component of IVIG, in particular, appears to be critical for these properties.^{6,7}

The beneficial pathobiochemical effects of IVIG have been previously described.^{8,9} The role of sepsis therapy using traditional IgG-enriched IVIGs for improving survival in severe sepsis and septic shock in humans remains controversial. Most clinical IVIG studies in humans are relatively small. One large, multicenter, randomized, controlled trial in adult patients ($n = 624$)¹⁰ and one in infants with neonatal sepsis ($n = 3493$)¹¹ found no survival benefit for patients treated with established IVIG solutions. IgM-enriched immunoglobulin solutions appear to have a survival benefit in humans.¹² There is a discrepancy between the experimental IVIG data^{5,8,9} and the clinical results.¹³ One possible factor might be the spectrum of immunoglobulin subtypes used in the IVIG solutions.

In this study, we assessed the ability of BT086, an IgM-enriched IVIG solution (23% of all immunoglobulins are IgM), to eliminate *Escherichia coli* in anesthetized rabbits with endotoxemia. We further assessed the effects of BT086 administration on bacterial distribution, residual granulocyte phagocytic activity, respiratory burst activity, histopathological changes in lung and liver tissues, and inflammatory mediator plasma levels in endotoxemic rabbits.

Materials and methods**Rabbit model of endotoxemia**

The techniques of preparing and conducting experiments in our rabbit endotoxemia model have previously been described in detail.^{14,15} All experiments were performed after approval by the commission for animal protection of the local government (AZ 24-9168.11-1/2011-1). Twenty adult New Zealand rabbits were anesthetized and mechanically ventilated via a respirator (Evita, Dräger, Lübeck, Germany) during the entire observation period (240 min). The systemic arterial pressure, rectal temperature, and electrocardiogram were continuously monitored and digitally recorded ("ViPaD", PCat Computer, Dresden, Germany).

Bacterial culture

A serum-resistant and non-hemolytic strain of *E. coli* (*E. coli* 018ac:K1:H7, ATCC 700973) with a smooth LPS phenotype was cultivated in tryptic soy broth and frozen in aliquots with glycerol at -80°C until use. Freshly grown *E. coli* cultures were prepared for the animal experiments. According to our pilot experiments, a standardized *E. coli* inoculum of 10^8 CFU was selected to enable the detection of cultivable bacteria in control animals until the end of the experiments.

Experimental protocol

The animals were randomly assigned to one of the following four groups ($n = 5$ per group): (1) the negative control group without LPS (control group), (2) the positive control group with LPS infusion (LPS group), (3) the albumin-treated LPS group (ALB + LPS group) to ascertain the protein-related effects, and (4) the BT086-treated LPS group (BT086 + LPS group).

Each experiment lasted 240 min after instrumentation of the rabbits (Fig. 1). Following a 30-min stabilization period (t_{-30}), intravenous LPS (Sigma, Deisenhofen, Germany) was administered continuously at a rate of 40 $\mu\text{g}/\text{kg}/\text{h}$ to all the rabbits, except the animals in the control group. At t_0 , a standardized amount of *E. coli* was intravenously injected into all of the animals. After 15 min (t_{15}), the control group and the LPS group continuously received a 2 ml/kg/h infusion of balanced electrolyte solution E153 (Serumwerk, Bernburg, Germany). The animals in the albumin treatment group (ALB + LPS group) received 2 ml/kg/h 5% albumin solution (Biotest, Dreieich, Germany), and the animals in the BT086 treatment group (BT086 + LPS group) received 2 ml/kg/h BT086 solution (Biotest), as did a group in a previous study.¹⁵ For the analysis of the arterial blood gases, leukocyte counts, hematocrit, hemoglobin concentrations, and bacterial blood clearance, arterial blood was aseptically drawn at the time points t_{-30} , t_0 , t_1 , t_{15} , t_{30} , t_{90} , t_{120} , and t_{180} . The polymorphonuclear neutrophil (PMN) oxidative burst, residual phagocytic

activity, and inflammatory mediator levels were determined in the arterial blood at t_{-30} , t_{30} , t_{90} , and t_{180} .

At the end of each experiment, the animals were euthanized. Subsequently, tissue samples from the liver, spleen, kidney, and left lung were removed under aseptic conditions for the bacterial cultures. One sample from a central region of the right fixed lung (adjacent to the main bronchus) and two from a peripheral-dependent and non-dependent region of the lung were embedded in paraffin, stained with hematoxylin–eosin, and cut into slices for the morphometric and histological analysis.

Determination of the bacterial load

The blood samples and homogenized organs were serially diluted in normal saline and plated onto blood agar plates. After incubation of the cultures at 37°C for 24 h, the respective *E. coli* colonies were counted. The final bacterial concentration was calculated as the number of CFU per milliliter of blood or as CFU per gram of tissue.

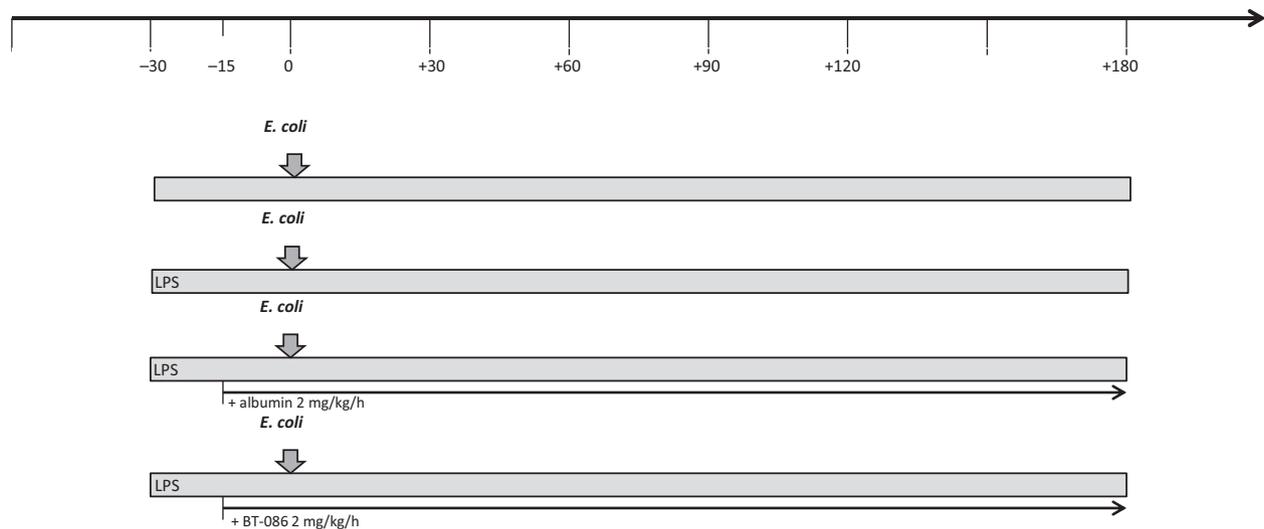


Fig. 1. Experimental setup. The animals were randomly assigned to one of the four groups: (1) the negative control group without LPS (control group), (2) the positive control group with LPS infusion (LPS group), (3) the albumin-treated LPS group (ALB+LPS group) to ascertain the protein-related effects, and (4) the BT086-treated LPS group (BT086 + LPS group). Following a 30-min stabilization period (t_{-30}), intravenous LPS was administered continuously at a rate of 40 $\mu\text{g}/\text{kg}/\text{h}$ to all the rabbits, except the animals in the control group. At t_0 , a standardized amount of *Escherichia coli* was intravenously injected into all of the animals. After 15 min (t_{15}), the control group and the LPS group continuously received a 2 ml/kg/h infusion of balanced electrolyte solution, the ALB + LPS group received 2 ml/kg/h 5% albumin solution, and BT086 + LPS group received 2 ml/kg/h BT086 solution.

Quantitative analysis of PMN phagocytosis and PMN burst activity

The phagocytosis test (Phagotest; Glycotop Biotechnology, Heidelberg, Germany) measures the percentage of phagocytes that have ingested FITC-labeled bacteria and their activity (the number of bacteria per PMN).¹⁶ The phagocytosis test was used to evaluate the residual *ex vivo* phagocytic activity of PMNs in whole blood after the animals already had received a bolus load of unlabeled *E. coli* at t_0 , as described in the experimental protocol. The obtained *ex vivo* values represent the idle PMN phagocytic capacity in our model, which is an inverse measure of the *in vivo* PMN phagocytic activity. Single-cell analysis was performed by flow cytometry (FACS; Becton Dickinson, Heidelberg, Germany). The residual PMN phagocytic capacity was determined by the intracellular content of FITC-marked *E. coli* and is expressed as the mean channel fluorescence per cell.^{15,16}

The extent of the PMN intracellular radical oxygen species production was determined using samples of freshly drawn heparinized blood. The oxidative burst of the PMNs was investigated using a test kit (Bursttest; Glycotop Biotechnology, Heidelberg, Germany). Single-cell analysis was performed by laser flow cytometry. The method of the quantitative assay for monitoring the oxidative burst has previously been described in detail.^{15,17}

Histopathological analysis

The histological samples were examined using digitalized photomicrographs of three non-overlapping fields from lung specimen. The images were digitalized and processed using a computer-based system and imaging analysis software (AnalySIS, version 3.1; Soft Imaging System, Münster, Germany). Diffuse alveolar damage (DAD) in the blinded samples was quantified systematically by the same investigator using a DAD scoring system. The pathological features of the lung in adult respiratory distress syndrome are characterized under the term DAD.¹⁸ The histopathological characteristics of the scoring system used in this study were adapted from the literature¹⁹ and modified

to include a weighing system similar to that proposed by Broccard et al.²⁰

For the morphometric analysis of the lung tissue, the images were binarized (the color images were converted to black and white images), with the black portions representing parenchyma, edema, or infiltration (non-aerated), and the white portions representing the aerated areas. The aerated and non-aerated areas were measured and appraised with imaging analysis software (AnalySIS, version 3.1; Soft Imaging System).

A periodic acid-Schiff stain was used for the detection of glycogen in the hepatocytes. The formalin-fixed, paraffin-embedded tissue sections were deparaffinized, hydrated and placed in Schiff reagent (SEV Liquid Solutions, Flintsbach a. Inn, Germany) with following counterstain with hematoxylin. For the quantitative analysis of the glycogen levels, the images were binarized and rated using image analysis software (AnalySIS, version 3.1; Soft Imaging System).

The protein levels of tumor necrosis factor- α (TNF α), interleukin-1 (IL-1), and interleukin-6 (IL-6) were measured in the blood plasma at time points t_{-30} , t_{30} , t_{90} , and t_{180} using commercial enzyme-linked immunosorbent assay kits (R&D Systems, Wiesbaden, Germany), according to the manufacturer's instructions.

Statistical analysis

The power analysis and previous studies using this model^{14,15} indicated that a caseload of five animals per group detects a 20% difference in the bacterial CFUs at the 180 min time point with a *P* value of less than 0.05 and a statistical power of 80%. The single measurements between the groups were compared with an unpaired two-sided Student's *t*-test. The between-group differences after the repeated measurements were tested by general linear model statistics, according to a two-way ANOVA. When applicable, the differences in the baseline values were determined by considering the baseline values as covariates within the statistical two-way ANCOVA. Significance was accepted at *P* < 0.05. The statistical analysis was performed with SPSS Statistics (release 20.0.0; IBM Corp., Armonk, NY, USA).

Results

Hemodynamic measurements and blood gas analysis

The measurements of the systolic, mean, and diastolic blood pressure, heart rate, rectal temperature, blood gases, and hemoglobin and hematocrit values as well as the differential blood counts and serum lactate levels during the time course of the experiment did not differ among the groups (data not shown).

Bacterial blood clearance and organ colonization

The elimination of *E. coli* from the bloodstream of the BT086-treated rabbits with endotoxemia was augmented compared with the untreated (LPS group, $P < 0.001$) and albumin-treated rabbits

with endotoxemia (ALB + LPS group, $P < 0.05$, Fig. 2). The first sterile blood culture was determined earlier in the BT086-treated rabbits with endotoxemia than in the untreated and albumin-treated animals with endotoxemia ($P < 0.05$, Fig. 3). No difference was detected between the control group (without LPS) and the BT086 + LPS group (Fig. 3). This finding suggests that the BT086 IgM-enriched solution improves the host defense by neutralizing the effects of LPS in bloodstream infections. The organ colonization did not differ between the groups.

Phagocytosis and burst activity of PMNs

The IgM effects on the host defense were analyzed by assessing the phagocytosis and burst capacity after the previous bloodstream inoculum. As described, the blood bacterial clearance

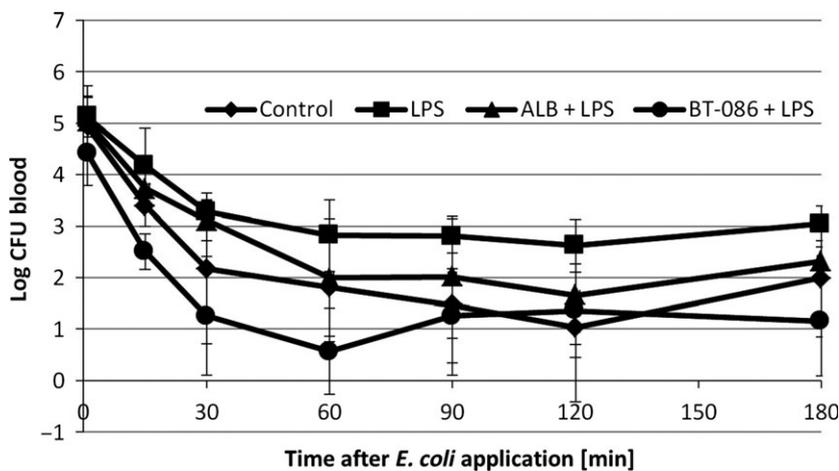


Fig. 2. *Escherichia coli* blood elimination kinetic. The blood clearance of *E. coli* (mean \pm SD) in the animals after injection of 10^8 CFU *E. coli*. Control: negative control group without LPS, LPS: positive control group with LPS infusion and no treatment, ALB + LPS: LPS group treated with albumin, BT086 + LPS: LPS group treated with BT086. LPS vs. control $P < 0.05$, LPS vs. BT086 $P = 0.001$

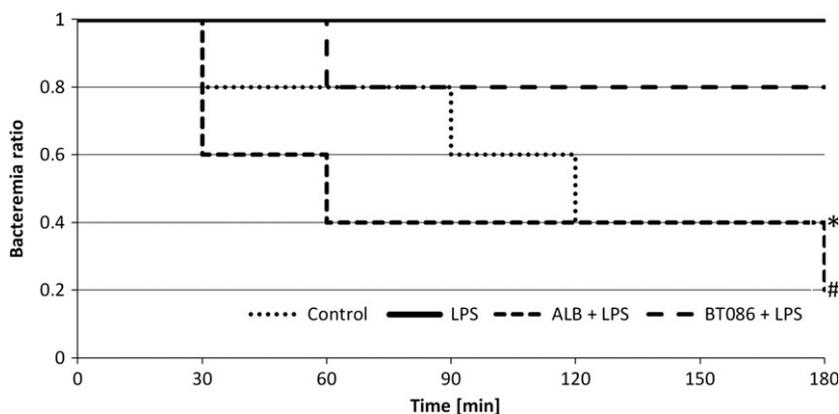


Fig. 3. Time to the detection of the first sterile blood culture. Kaplan-Meier estimate. Control group: negative control group without LPS, LPS: a positive control group with LPS infusion, ALB + LPS: LPS group treated with albumin BT086 + LPS: LPS group treated with BT086. * $P < 0.05$ LPS vs. control group, # $P < 0.05$ BT086 + LPS vs. LPS group

was assessed by measuring the inoculum of unlabeled *E. coli* during the *in vivo* phase of the experiment. At 180 min, the percentage of the additional phagocytic PMNs, indicated by the additional *ex vivo* uptake of FITC-marked *E. coli*, was lower in the BT086 + LPS group than in the LPS and ALB+LPS groups ($P < 0.05$, Fig. 4). At 180 min, the phagocytic activity, characterized by the number of phagocytized bacteria per leukocyte, was higher in the BT086 + LPS group ($P < 0.05$, Fig. 5). The oxidative burst of the PMNs did not differ in any of the groups.

Histopathological analysis

The DAD score analysis as well the morphometric analysis of the lung tissue did not differ between the groups (Fig. 6). The percentage of glycogen was higher in the BT086 + LPS group than in the other groups ($P < 0.001$, Figs. 7 and 8).

Inflammatory Mediators

The measurements of the TNF α , IL-1, and IL-6 plasma levels did not differ among the groups (Table 1).

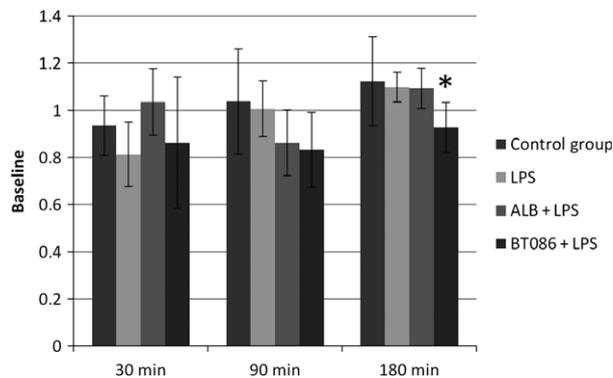


Fig. 4. Polymorphonuclear neutrophil residual phagocytosis. The percentage of additionally phagocytic PMNs. The percentage of additional phagocytic PMNs, indicated by the additional uptake of FITC-marked *Escherichia coli*, as a percent of the baseline measurements (t_{-30} min). At 180 min, the percentage of additional phagocytic PMNs was lower in the BT086 + LPS group compared with the LPS and ALB + LPS groups ($*P < 0.05$). All data are the mean \pm SD

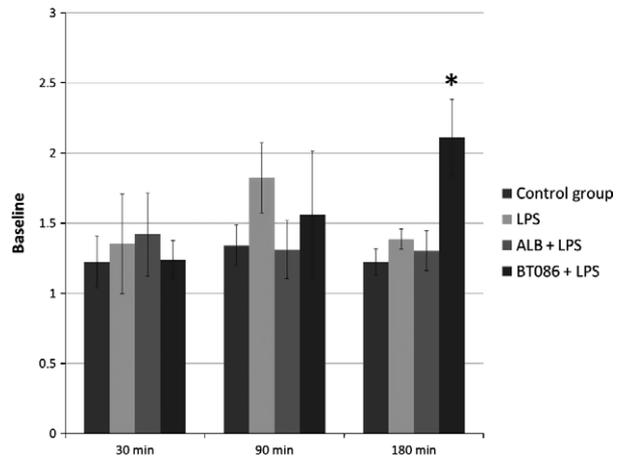


Fig. 5. Polymorphonuclear neutrophil residual phagocytosis. Phagocytes activity. The activity of the phagocytes, characterized by the number of phagocytized FITC-marked bacteria per leukocyte, as a percentage of the baseline measurements (t_{-30} min). At 180 min, the phagocytic activity was higher in the BT086 + LPS group ($*P < 0.05$). All data are the mean \pm SD.

Discussion

In this study, we present the first experimental data using BT086, a modified IgM-enriched solution, in an established animal model of experimental endotoxemia and demonstrate that BT086 improved the host defense and preserved the energy stores in the liver in experimental endotoxemia in rabbits.

The potential use of IgG-enriched IVIG solutions to combat bacterial endotoxin and inflammatory dysregulation in sepsis has been of major interest in recent decades. Despite the promising effects of human IgG-enriched IVIG solutions in animal experiments,²¹ conventional class G-based immunoglobulin therapy did not improve clinical outcome of critically ill patients with sepsis.²² The spectrum of the immunoglobulin subtypes used in the IVIG solutions might be important to the clinical effect. Two meta-analyses demonstrated a trend toward superior results with IgM-enriched solutions compared with those enriched with IgG.^{12,23} The rationale for enrichment of IVIGs with IgM-type immunoglobulins is the unique ability of IgM to neutralize LPS in the early stage of antibody-dependent host defense.¹² Specifically, IgM contributes to the opsonization of bacteria,⁷ inhibits the LPS-triggered release of TNF α in mono-

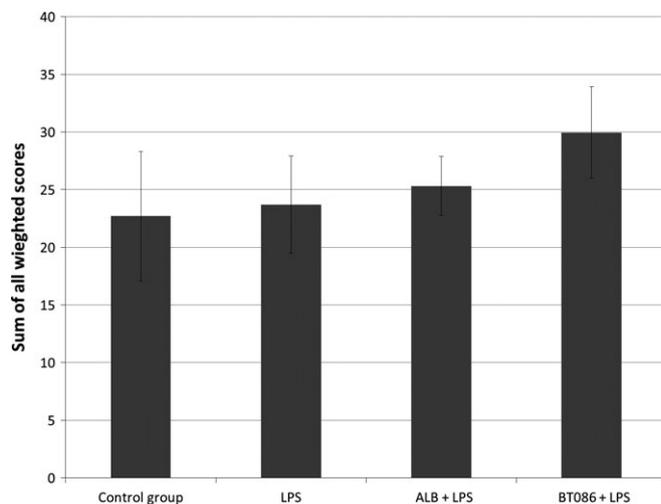


Fig. 6. The sum of all weighted diffuse alveolar damage scores. The sum of all of the weighted diffuse alveolar damage scores showed no difference between the groups. All data are the mean \pm SD.

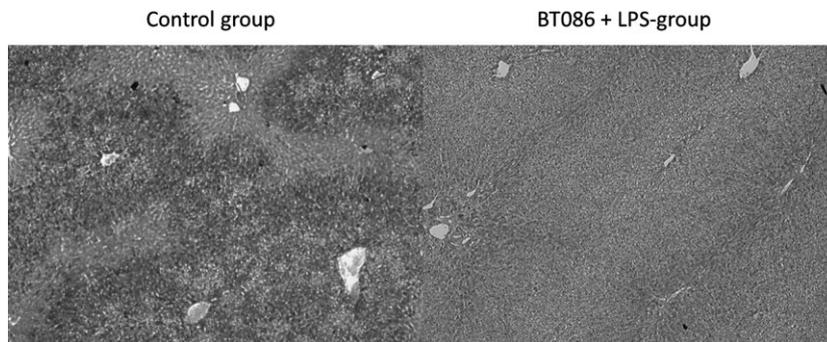


Fig. 7. Periodic acid-Schiff stain of the hepatocytes. The periodic acid-Schiff stains demonstrate the glycogen content of the hepatocytes. The glycogen content of the hepatic cells from the BT086 + LPS group (right) is considerably higher compared to all other groups (the control group is shown on the left).

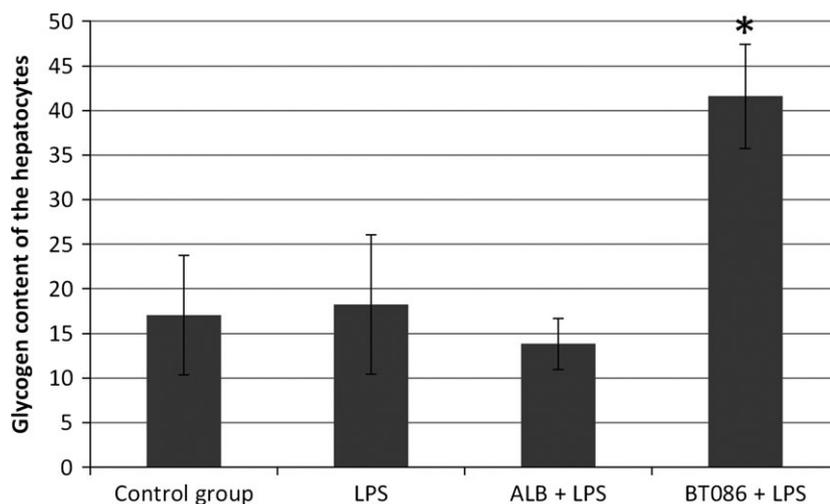


Fig. 8. The glycogen content of hepatocytes. The periodic acid-Schiff stain of the hepatocytes in the BT086 + LPS group showed a significantly higher glycogen content compared to the control group and the ALB + LPS group (* $P < 0.001$). All data are the mean \pm SD.

cytes,⁶ and stimulates the production of antibodies against LPS,⁷ which is one of the key activators of the complex anti-inflammatory cascade.^{12,24}

In this study, we evaluated an IVIG solution with an increased portion of IgM. Using the present model of endotoxemia in rabbits, our group previously examined the effects of IgG²¹- and

Table 1 The plasma level of TNF α , IL-1, and IL-6 before and after the induction of endotoxemia in anesthetized rabbits.

| | Control | LPS | ALB | BT086 |
|----------------------|------------------|-----------------|------------------|------------------|
| TNF α , pg/ml | | | | |
| -30 min | 87 \pm 196 | 0 | 0 | 0 |
| +30 min | 640 \pm 994 | 1934 \pm 2043 | 1449 \pm 467 | 632 \pm 384 |
| +90 min | 7179 \pm 2080 | 4766 \pm 3668 | 6913 \pm 4046 | 5289 \pm 6561 |
| +180 min | 61 \pm 36 | 70 \pm 91 | 118 \pm 123 | 331 \pm 564 |
| IL-1, pg/ml | | | | |
| -30 min | 82.0 \pm 25.6 | 68.0 \pm 37.4 | 79.0 \pm 35.5 | 95.3 \pm 12.8 |
| +30 min | 80.7 \pm 30.6 | 70.3 \pm 36.8 | 74.4 \pm 31.4 | 93.8 \pm 5.3 |
| +90 min | 91.5 \pm 20.8 | 87.8 \pm 41.9 | 88.4 \pm 28.2 | 114.6 \pm 6.7 |
| +180 min | 110.2 \pm 26.7 | 91.9 \pm 37.1 | 100.5 \pm 27.1 | 123.5 \pm 16.2 |
| IL-6, pg/ml | | | | |
| -30 min | 80.0 \pm 34.2 | 56.3 \pm 35.1 | 68.5 \pm 20.5 | 97.4 \pm 38.6 |
| +30 min | 78.8 \pm 51.2 | 60.6 \pm 32.9 | 75.4 \pm 34.6 | 91.0 \pm 26.5 |
| +90 min | 91.4 \pm 39.9 | 79.7 \pm 45.6 | 88.8 \pm 28.0 | 102.5 \pm 26.8 |
| +180 min | 101.2 \pm 46.9 | 83.5 \pm 41.3 | 95.6 \pm 42.8 | 115.6 \pm 45.7 |

The measurements of TNF α , IL-1, and IL-6 levels did not differ among the groups (all the data are the mean \pm SD).

IgM-enriched solutions.¹⁵ The *E. coli* dose was determined in a pilot study to ensure that bacteremia could be detected in healthy animals until the end of the experiments. Additionally, we continuously administered intravenous LPS in the previously identified dose^{14,25} in all the groups except the control. We achieved endotoxemia with bacteremia, allowing for the detection of the *in vivo* effects on early phase of bacterial clearance. To control for non-specific protein-related effects, we established an albumin-treated LPS group (ALB + LPS group). In this study, the time to the first sterile blood culture was shorter in the BT086 + LPS group than in the LPS and ALB + LPS groups, without a significant difference compared to the control group (Fig. 3). Treatment with BT086 appeared to neutralize the LPS effect concerning bacterial clearance. Previous findings by our group showed no significant effects on the bacterial blood elimination kinetics in animals receiving an IgM-enriched solution¹⁵ possibly due to another composition of the immunoglobulins in the BT086 compared to Pentaglobin[®], which was used in earlier studies. In the previous study, the examination of the IgM effects on the *E. coli* organ concentrations showed that IgM reduced the *E. coli* counts in the liver and spleen of rabbits. This observation was interpreted as an improved reticuloendothelial system clearance effect of the IgM-enriched solution.¹⁵

In the present study, we found no differences in the organ colonization of *E. coli*, which is most likely a consequence of the enhanced bacterial blood clearance. Our findings are in agreement with those of Fabrizio et al., who found that a single dose of IgM led to a reduction in the bacterial blood load.²⁶ As a possible explanation, the authors discussed a promotion of *in vivo* killing. Phagocytosis by competent cells and the production of reactive oxygen species play a crucial role in the innate immune response to eliminate the invading microorganism.²⁷ We found a significantly higher phagocytic activity in the BT086 + LPS group at the end of experiment, without a difference in the oxidative burst of PMNs in any group. The higher number of phagocytized bacteria per leukocyte might be one of the reasons for the improved bacterial clearance observed in the BT086 + LPS group. These data agree with our previous study,¹⁵ in which we demonstrated an enhanced *in vivo* phagocytosis as depicted in electron microscopy. Gille et al. reported similar results in experiments using human cord blood and peripheral blood monocytes.²⁸ The authors found that the addition of immunoglobulins to freshly harvested monocytes increased their phagocytic capacity, measured via the number of phagocytic monocytes and the amount of bacteria per cell. In agreement with our results, Tinguely et al. found that excess immunoglobulin preparations

promoted *in vitro* phagocytosis, more so for monocytes than for PMNs alone.²⁹ Danikas et al. showed that the phagocytic activity of peripheral blood PMNs in severe sepsis patients at the time of admission was lower in the patients who later died.³⁰ This finding is in agreement with the well-described fact that the immune system becomes hyporeactive during the later stages of sepsis, which is characterized by several hallmarks including increased anti-inflammatory cytokine release³¹ and ablated blood leukocyte responsiveness.^{32,33} It appears that BT086 might hamper the immunosuppression of peripheral blood PMNs or increase their activity because our endotoxemia and sepsis model demonstrated enhanced phagocytosis after the administration of BT086. This finding highlights the possible clinical benefit of BT086 for septic patients, at least in the early phase of sepsis.

We found no differences in the plasma levels of inflammatory mediators such as TNF α , IL-1, and IL-6 between the groups. Some authors have described the anti-inflammatory effects of intravenous immunoglobulins such as a neutralizing effect on microbial toxins and pro-inflammatory cytokines,³⁴ the inactivation of the potent pro-inflammatory anaphylatoxins, C3a and C5a,³⁵ and significantly reduced levels of keratinocyte-derived chemokine, IL-6, and macrophage inflammatory protein-2 expression.²⁶ Barrat-Due et al. documented early inflammatory responses to Pentaglobin in a porcine model of *E. coli*-induced sepsis.³⁶ In our study, neither a pro-inflammatory nor an anti-inflammatory effect of the selected dose of was discernible, based on the evaluation of the inflammatory mediators, the diffuse alveolar damage, and the morphometric lung tissue analysis. We found no effects of BT086 on the hemodynamic responses. Possibly, the absence of significant effects of BT086 treatment on the hemodynamic parameters and inflammation is a consequence of the relatively short observation period in our model of endotoxemia and early sepsis. In addition, the failure to demonstrate any difference between groups on inflammatory markers could potentially be a relatively small group size and large SD on cytokine data.

In humans, glycogen in the muscles and the liver represents one form of energy storage. During sepsis, patients frequently enter a

hypermetabolic state, marked by several alterations including increased glucose production and depressed gluconeogenesis,³⁷ with consequential depletion of the glycogen stores. To our knowledge, no data exist on the effects of immunoglobulins on the glycogen content in hepatocytes. We demonstrated a significantly higher concentration of glycogen in hepatic cells after BT086 treatment, as opposed to that in the other groups. The clinical importance of this finding remains unclear; however, it is possible that the increased energy stores in the form of glycogen have a beneficial effect during critical illness.

In conclusion, in this study we demonstrate improvements in bacterial blood clearance and phagocytic activity, as well a higher glycogen content in hepatocytes, after application of the IgM-enriched solution, BT086 (Biotest, Dreieich, Germany). Our data suggest that IgM-enrichment of IVIGs has the potential to improve the host defense by neutralizing the effects of LPS on bloodstream infection in a rabbit model of endotoxemia and sepsis. Additional animal studies using different models of sepsis and inflammation, as well as using a different dosage of BT086, are necessary to further explore the potential beneficial effects of BT086. With these findings, a positive influence of IgM-based immunoglobulin therapy on host defense in endotoxemia could be suggested.

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