

Lactoferrin reduces febrile neutropenia in children receiving chemotherapy for hematologic malignancies: a randomized, placebo-controlled trial

Febrile neutropenia (FN) is one of the most frequent and clinically significant complications of intensive chemotherapy for pediatric hematologic malignancies, occurring in up to 25% of cases.¹ FN can lead to bloodstream infection, septic shock, and treatment delays. In addition to its direct morbidity, FN results in prolonged hospitalization, increased healthcare costs, and overuse of antibiotics.² Notably, exposure to systemic antibiotics during FN episodes increases the risk of microbial resistance, antibiotic-related adverse events, and disruption of the gut microbiota (GM).³ Recent evidence indicates that the GM also plays a pivotal role in the pathogenesis of FN. Chemotherapy-induced mucosal injury leads to a loss of microbial diversity and an overgrowth of pathobionts, such as *Enterobacteriaceae*, *Enterococcaceae*, and *Akkermansia spp.*, which can facilitate bacterial translocation and trigger systemic inflammation.⁴ Therefore, strategies that preserve a healthy microbiota may offer an effective, antibiotic-sparing approach to infection prevention.

Lactoferrin (Lf) is an 80 kDa cationic glycoprotein secreted by epithelial cells and neutrophil granules and is abundant in colostrum and other mucosal secretions. Lf exhibits antimicrobial, anti-inflammatory, and immunomodulatory properties, acting through mechanisms such as iron sequestration, disruption of bacterial membranes, inhibition of biofilm formation, and modulation of cytokine release (including interleukin [IL]-6, IL-10, and tumor necrosis factor α [TNF- α]).^{5,6} Notably, oral bovine Lf (bLf), which shares 70% sequence homology with the human protein, has demonstrated clinical benefits in reducing sepsis and necrotizing enterocolitis in preterm infants^{7,8} as well as in mitigating infectious diarrhea and recurrent respiratory or urinary infections in children.⁹

Given these properties, we hypothesized that oral bLf could reduce the incidence of FN and infections during induction chemotherapy in children with acute lymphoblastic leukemia (ALL), acute myeloid leukemia (AML), or non-Hodgkin lymphoma (NHL). These diseases are all characterized by profound and prolonged neutropenia, mucositis, and gut microbiota dysbiosis.⁴

We conducted a prospective, multicenter, randomized, double-blind, placebo-controlled trial (*clinicaltrials.gov*. Identifier: NCT07113314) among nine centers of the Italian Association of Pediatric Hematology and Oncology (AIEOP). The study was approved by the Ethics Committee of Fondazione IRCCS Policlinico San Matteo, Pavia, Italy (protocol 20160001176), and ratified by local ethics boards.

Eligible patients were aged >1 month and <21 years, receiving first-line induction chemotherapy for ALL, AML, or NHL according to AIEOP protocols, with written informed consent signed from parents or guardians. Exclusion criteria included gut colonization with multidrug-resistant organisms, prior chemotherapy, or inability/refusal to take oral medication.

Patients were randomized 1:1, stratified by center, using blocks of different size (2-6). Random allocation sequence was generated using the software STATA 14.0 (StataCorp., College Station, TX, USA). Allocation was managed by an Interactive Web Response System (REDCap-based).

The experimental arm received oral bLf (Mosiatic®, Pharmaguida, Rome, Italy) 200 mg once daily for 8 weeks, corresponding to the induction chemotherapy period. The control arm received identical placebo capsules. Both products were indistinguishable in appearance, ensuring that study staff, patients, and families remained blinded to group allocation. Capsules can be opened and the powder swallowed with few mL of liquids. Adherence was monitored through patient diaries and capsule counts. The purity of bLf, checked by SDS-PAGE and silver nitrate staining, was about 98%. The bLf iron saturation was about 10%. All supportive therapies were according to local practices, and the use of antibiotic prophylaxis was recorded in the electronic CRF.

The primary endpoint was the incidence of FN with absolute neutrophil count (ANC) <0.1 g/L over 90 days. Secondary endpoints included incidence of FN with ANC <0.5 g/L, fever of unknown origin (FUO), microbiologically documented infections, and safety.

This preliminary efficacy study aimed to exclude, using a one-sided 80% confidence interval (CI), that the difference in FN incidence between the treated and control groups was less than 5%.

Patients were analyzed according to their randomized treatment group, using intention-to-treat analysis. Incidence rates were compared by Cox regression and Kaplan-Meier analysis, with results expressed as hazard ratios (HR). The number needed to treat (NNT) was calculated as the inverse of cumulative incidences difference between the two treatment arms at specific points in time derived from the Kaplan-Meier curves.

Between January 2017 and June 2020, 156 patients were enrolled: 79 in the bLf arm and 77 in the placebo arm, respectively (*Online Supplementary Figure S1*). Median age was 6 years (range, 1-17) in the bLf group and 7 years

(range, 1-20) in the placebo group. Diagnoses included ALL (117/156, 75.0%), AML (17/156, 10.9%), and NHL (22/156, 14.1%). At enrollment, there were no between-arm differences in clinical and demographic data (Table 1).

Over 90 days, FN with ANC <0.1 g/L occurred in 33 of 79 patients (41.8%) in the bLf arm versus 40 of 77 (51.9%) in placebo. Accounting for time-to-event in a Cox regression analysis, the incidence rate was significantly lower in the bLf arm (4.4 vs. 6.9 cases per 100 person-weeks), corresponding to a 37% risk reduction (HR=0.63; 80% CI: 0.47-0.85; P=0.05) (Figure 1). The NNT at 90 days was eight, meaning one FN event was prevented for every eight patients treated. The NNT at weeks 4, 6, and 8 were seven, eight and nine, respectively.

When considering FN with ANC <0.5 g/L, the incidence was still lower in the bLf group (58.2%) compared to placebo (63.6%), and in a Cox regression model the incidence rate of this event tended (P=0.10) to be lower (-29%) in the bLf group (7.1 cases per 100 persons-week) as compared to the placebo group (9.9 cases per 100 persons-week) (bLf vs. placebo; HR=0.71; 80% CI: 0.55-0.93) (Online Supplementary Figure S2). At 4, 6, and 8 weeks, the NNT values for the incidence of FN <0.5 g/L were ten, ten, and 11, respectively. During the 90-day follow-up, FUO occurred in 36 of 77 (46.7%) placebo versus 32 of 79 (40.5%) bLf patients - 19% lower total number of episodes (43 vs. 53). The 90-day cumulative incidence was 50% (95% CI: 38-62) in placebo and 42% (95% CI: 31-53) in bLf (P=0.20; NNT=12) (Figure 2A). MDI were diagnosed in 11 of 77 (14.3%) placebo and eight of 79 (10.1%) bLf patients (P= 0.40) (Figure 2B). The distribution of pathogens among blood cultures was comparable between groups (Online Supplementary Table S1). Incidence of *Clostridioides difficile*, vancomycin-resistant *Enterococcus*, or multidrug-resistant bacteria was too low for analysis. No treatment-related adverse events (AE)

were observed; overall AE incidence was not statistically different between groups (placebo arm, 18 AE in 14 patients; bLf arm, 26 AE in 10 patients).

This is the first randomized, double-blind, placebo-controlled trial providing evidence that oral bLf may reduce FN incidence during induction chemotherapy in children with hematologic malignancies. Patients receiving bLf experienced a 37% lower rate of FN and a consistent trend toward fewer FUO episodes, without safety concerns. To our knowledge, this is the first randomized trial evaluating bLf as a preventive strategy for FN in pediatric oncology. The effect size observed exceeded initial expectations and aligns with prior evidence of LF's protective role against sepsis in preterm infants.¹⁰

Lf's antimicrobial actions are multifaceted. Through iron sequestration, it deprives pathogens of an essential growth

Table 1. Patients' characteristics by study arms.

Characteristics	Patients		Total
	Placebo group N=77	bLF group N=79	N=156
Sex, N (%)			
Male	42 (54.6)	49 (62.0)	91 (58.3)
Female	35 (45.5)	30 (38.0)	65 (41.7)
Age at random, years, median (min-max)	7 (1-20)	6 (1-17)	6 (1.0-17)
Underlying diseases, N (%)			
ALL	56 (72.7)	61 (77.2)	117 (75.0)
AML	9 (11.7)	8 (10.1)	17 (10.9)
NHL	12 (15.6)	10 (12.6)	22 (14.1)

ALL: acute lymphoblastic leukemia; AML: acute myeloid leukemia; bLF: bovine lactoferrin; NHL: non-Hodgkin lymphoma; min: minimum; max: maximum.

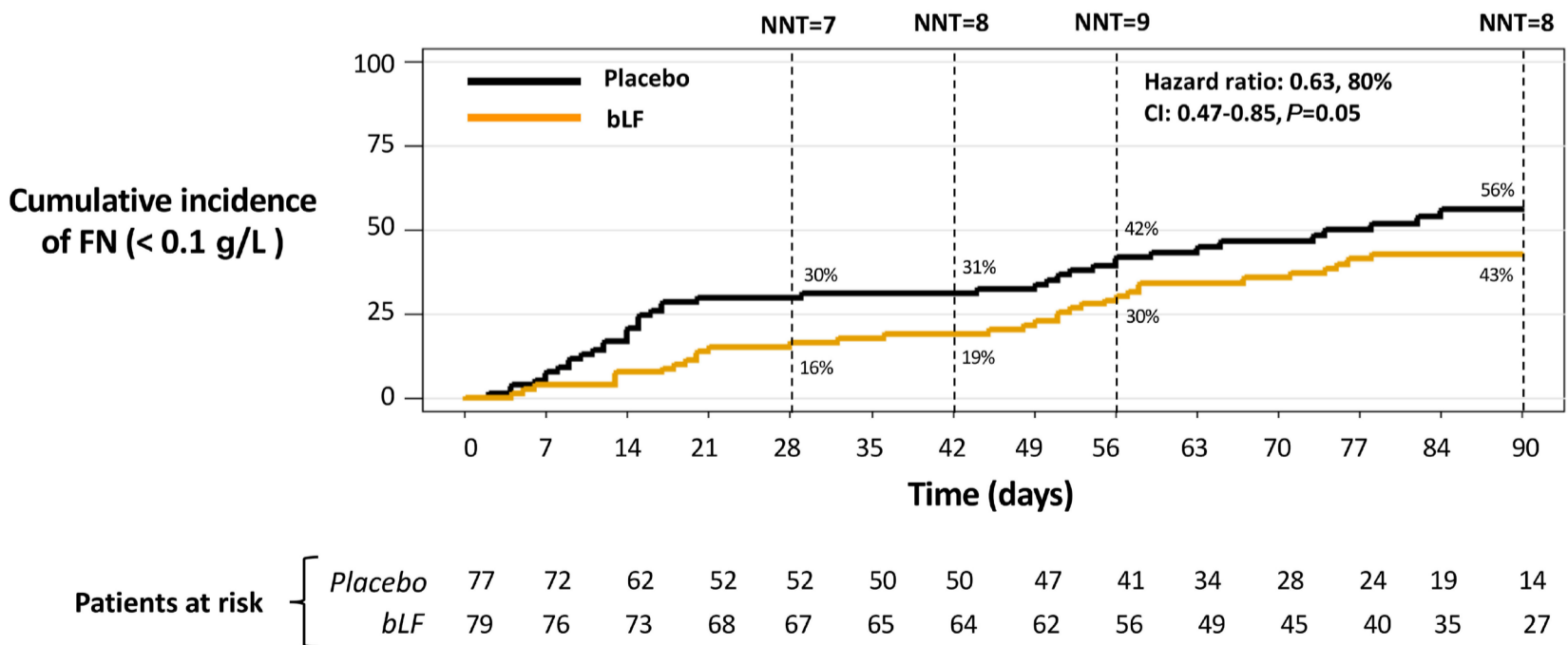


Figure 1. Cumulative incidence of febrile neutropenia by study arms. bLF: bovine lactoferrin; CI: confidence interval; FN: febrile neutropenia; NNT: number needed to treat.

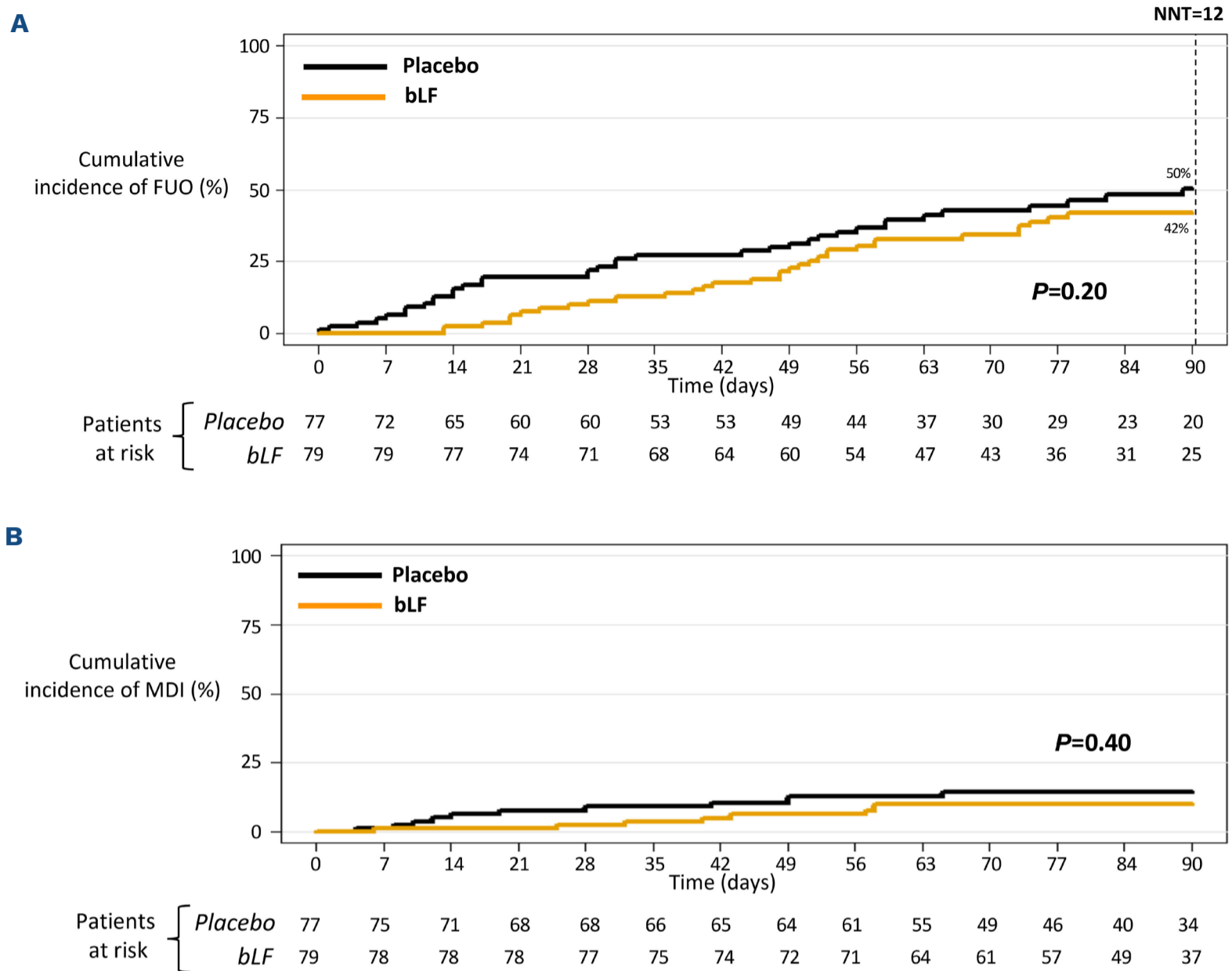


Figure 2. Cumulative Incidence of fever of unknown origin and microbiologically documented infection by study arms. (A) The cumulative incidence of fever of unknown origin (FUO) across study arms. (B) The cumulative incidence of microbiologically documented infection (MDI) in the 2 groups. bLF: bovine lactoferrin; NNT: number needed to treat.

factor, limiting proliferation of Gram-positive and Gram-negative bacteria and fungi. In addition, Lf binds bacterial lipopolysaccharides and lipoteichoic acid, disrupts membranes, inhibits adherence and internalization into host cells.¹¹ Beyond direct antimicrobial effects, Lf exerts anti-inflammatory and immunomodulatory actions. It modulates cytokine production (IL-1 β , IL-6, TNF- α), promotes IL-12-mediated Th1 responses, enhances natural killer and T-cell activity, and reduces oxidative stress via iron chelation.¹² These effects may help explain why, in our cohort, the incidence of microbiologically documented infections and bacteremia was similar between groups, whereas the incidence of FUO was reduced in bLf-treated patients - particularly during the first 6 weeks of treatment. This pattern is reflected in the Kaplan-Meier cumulative-incidence curve, which mirrors the trend observed for FN. The origin of FUO is often non-infectious and may be driven by cytokine-mediated mechanisms. Another mechanism may involve preservation of healthy gut microbiota, as found in an ancillary study of our cohort,¹³ where bLf prevented chemotherapy-induced loss

of microbial diversity, reduced pathobionts (*Enterobacteriaceae*, *Akkermansia*), and maintained butyrate-producing commensals. These changes are consistent with prior work showing that *Enterococcus* and *Akkermansia* expansion predicts FN and infection risk in leukemia patients.⁴ Maintenance of butyrate producers supports mucosal integrity, modulates inflammatory responses, and decreases oxidative stress, potentially mediating the clinical benefit of bLf. Future studies should explore extended supplementation throughout intensive chemotherapy, weight-adjusted dosing, and evaluation in higher-risk settings such as relapsed leukemia or hematopoietic stem cell transplantation. Combining bLf with other microbiota-modulating interventions (e.g., prebiotics or postbiotics) may further enhance efficacy.¹⁵ In conclusion, oral bLf supplementation during induction chemotherapy may reduce FN incidence in children with acute leukemia or lymphoma, without adverse effects. By bridging antimicrobial, anti-inflammatory, and microbiota-stabilizing mechanisms, bLf may represent a promising,

safe, and non-antibiotic strategy for preventing FN in high-risk pediatric patients. These encouraging findings warrant confirmation in a larger, adequately powered phase III trial.

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Disclosures

No conflicts of interest to disclose.

Contributions

ND developed the concept, performed the study, analyzed data and wrote the manuscript. DZ performed the study, analyzed data and wrote the manuscript. EM analyzed data, took care of patients and helped writing the manuscript. GT, CK and LS performed the biostatistical analyses, created the figures, and helped writing the manuscript. PM, KP, VV, AC, NG, RM, RDS and SR performed the study and took care for patients. MZ and SC critically revised the manuscript and analyzed data. All authors have critically revised and approved the final version of the manuscript.

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Data-sharing statement

For original data, please contact the corresponding authors.

Individual participant data will not be shared.

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