Evidence of pediatric sepsis caused by a drug resistant *Lactococcus garvieae* contaminated platelet concentrate

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**ABSTRACT**

Owing to an increasing number of infections in adults, *Lactococcus (L.) garvieae* has gained recognition as an emerging human pathogen, causing bacteraemia and septicemia. In September 2020, four paediatric onco-hematologic patients received a platelet concentrate from the same adult donor at Bambino Gesù Children’s Hospital IRCCS, Rome. Three of four patients experienced *L. garvieae* sepsis one day after transfusion. The *L. garvieae* pediatric isolates and the donor’s platelet concentrates were retrospectively collected for whole-genome sequencing and shot-gun metagenomics, respectively (Illumina HiSeq). By de novo assembly of the *L. garvieae* genomes, we found that all three pediatric isolates shared a 99.9% identity and were characterized by 440 common SNPs. Plasmid pUC11C (confering virulence properties) and the temperate prophage Plg-Tb25 were detected in all three strains. Core SNP genome-based maximum likelihood and Bayesian trees confirmed their phylogenetic common origin and revealed their relationship with *L. garvieae* strains affecting cows and humans (bootstrap values >100 and posterior probabilities = 1.00). Bacterial reads obtained by the donor’s platelet concentrate have been profiled with MetaPhlAn2 (v.2.7.5); among these, 29.9% belonged to Firmicutes, and 5.16% to Streptococcaceae (>97% identity with *L. garvieae*), confirming the presence of *L. garvieae* in the platelet concentrate transfusion. These data showed three episodes of sepsis for the first time due to a transfusion-associated transmission of *L. garvieae* in three pediatric hospitalized hematology patients. This highlights the importance to implement the screening of platelet components with new human-defined pathogens for ensuring the safety of blood supply, and more broadly, for the surveillance of emerging pathogens.

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**Inception**

*Lactococci* are Gram-positive, catalase-negative, facultative anaerobic cocci in short chains or pairs traditionally considered to be of low virulence to human beings [1]. Among them, *Lactococcus (L.) garvieae* species was first described in the 1950s in Japan, when it was discovered in a rainbow trout farm [2,3]. It can be found in a vast variety of environments due to its ability to adapt easily. It has been isolated from aquaculture, rivers, and sewage waters. The host range of *L. garvieae* is not limited to aquatic species, but is also associated with subclinical mastitis in cows and water buffalos, and pneumonia in pigs [4,5]. Previous studies have reported an association between *L. garvieae* infection and contaminated food, such as raw milk, cheese, vegetables, cereals, and meat [6,7].

The first case of *L. garvieae* human infection was observed in 1991 [8]. Since then, the relevance and clinical significance in humans have increased. After the first documented case, more than 30 new cases of *L. garvieae* infection have been described in literature [6,9–11]. Due to the increasing number of human clinical infections, *L. garvieae* has gained recognition as an emerging human pathogen. Among the reported cases, majority have been associated with bacteraemia with infective endocarditis among elderly and immunocompromised patients [9]. Other clinical syndromes include spontaneous septicemia,
liver abscess, bone infections, diverticulitis, and secondary peritonitis [3,9,11]. It has also been observed to mainly cause urinary tract infection (UTI) [1]. However, the true incidence of the disease is difficult to assess because of the morphological and biochemical similarities to other Gram-positive cocci like Enterococcus spp. and Streptococci [9,12].

In 2018, the first case of transfusion-transmitted L. garvieae resulting from a platelet transfusion in an adult hospitalized individual was described [13]. Here, we describe a cluster of transmission, with three cases of sepsis in onco-hematologic pediatric patients, derived by transfusion-transmitted L. garvieae from platelet concentrates (PCs) of the same donor.

**Material and methods**

**Study population**

On 31 August 2020, blood donation was collected from a healthy 59-year-old Italian female at IRCCS Bambino Gesù Children’s Hospital, in Rome, Italy. PCs were obtained by apheresis collection. The PCs were confirmed to have no abnormalities by visual inspection at the blood centre and again at the bedside just before starting the transfusion. These PCs were transfused to four paediatric patients at different times: patient 1 was transfused one day after blood donor’s collection, while patients 2–4 were transfused on day 2 after blood collection (hour:minute: second, respectively: 14:04, 14:32, and 16:36, respectively).

**Data and samples collection**

Age, sex, data regarding antibiotic therapies, comorbidities, and clinical conditions (body temperature, laboratory tests, and vital parameters) on the day of transfusion were collected for each patient.

At least one blood culture (one bottle for aerobic growth and one for anaerobic growth) was drawn for each patient before the transfusion as part of routine care. The Microbiology Unit of the Hospital processed the samples. According to the standard laboratory operating procedures, the identification of microorganisms was performed by MALDI-TOF mass spectrometry. As no breakpoints for antibiotic susceptibility have been determined for Lactococcus spp., antibiotic susceptibility was analyzed by the Vitek system according to the EUCAST guidelines for Streptococci. Antibiotic susceptibility was analyzed against the glycopeptides, teicoplanin and vancomycin; the penicillins, ampicillin and penicillin G; the cephalosporin, ceftriaxone; the lincosamide, clindamycin; the macrolide, erythromycin; the aminoglycoside, gentamycin; and the fluoroquinolone, levofloxacin.

Sepsis disease is defined as a suspected or proven infection with at least two of the following criteria: abnormal white-blood-cell count (elevated [>20,000 × 10⁹ per L] or decreased [<4000 × 10⁹ per L] for age), tachycardia or bradycardia, or tachypnoea [14]. C-reactive protein greater than 15 mg/L, and serum procalcitonin increase above 0.05 ng/mL [15].

**Ethics approval**

This study was conducted with respect to the Helsinki Declaration, and all the participants (parents) signed an informed consent to allow the use of clinical data for research purposes. Ethical approval was obtained from the Ethics Committee of Bambino Gesù Children’s Hospital in Rome, Italy (reference ID 2602/2021).

**Whole-genome sequencing and metagenomic analysis of donor’s platelet concentrates**

**Extraction and sequencing**

Microbial DNA was extracted from the positive cultures and the donor’s blood samples using the QIAamp DNA Microbiome Kit (Qiagen, Germany) according to the manufacturer’s instructions. DNA was quantified using the Qubit fluorometer. Libraries were obtained by QIAseq FX Single Cell DNA Library kit and DNAs were paired-end sequenced (2 × 150 bp) using Illumina HiSeq (Illumina, USA). All bacterial sequence data obtained were screened for the evidence of baseline quality control for short and long sequences. The quality check was done with FastQC v0.72.14. Adaptors were clipped and quality trimmed using FASTP 0.20.1 using default parameters (Q > 30). Quality trimmed paired reads were assembled into contigs using BYSS v2.0, a de-novo assembler based on de Brujin graph path reconstruction [16]. K-mer size optimization has been reached by comparing QUAST indices [17] obtained by assembling bacterial genomes with different k-mer sizes and by selecting the one returning the best score.

**Phylogenetic analysis**

To explore a possible clonal origin of the three L. garvieae strains isolated from paediatric patients, the L. garvieae genomes were compared to 20 publicly available L. garvieae sequences by both maximum likelihood and Bayesian approaches. Four publicly available additional Lactococcus (L.) lactis sequences were used as an outgroup. The characteristics of the 24 reference genomes are described in Table S1. PhaME software was used to perform the core single nucleotide polymorphism (SNP) genome typing [18]. The core SNP alignment was indeed composed of 27 sequences 62,990 nucleotide long. Phylogenetic relatedness was first analyzed by the maximum likelihood (ML) method using iqTree2 [19], under the nucleotide substitution GTR+I+G4 model [20] and 1000
bootstrap replicates. The results were confirmed by a Bayesian inference analysis through BEAST v.1.10.4 [21] by setting a chain length of 100 million of states under a strict molecular clock model and the GTR+I +G4 substitution model. The core SNP multiple alignment composed of the three L. garvieae strains isolated from paediatric patients plus 24 publicly available L. garvieae and L. lactis strains used for phylogenetic analysis is available at doi: 10.5281/zenodo.6473676. To better appreciate the conserved residues between Enterococcus, Streptococcus, and Lactococcus species, the core SNP multiple alignments composed of (i) the three L. garvieae strains isolated from paediatric patients plus 24 publicly available L. garvieae, L. lactis, and Streptococcus (S.) pneumoniae strains, (ii) the three L. garvieae strains isolated from paediatric patients plus three publicly available L. lactis, S. canis, and Enterococcus (E.) raffinosus strains are available at the same doi: 10.5281/zenodo.6473676.

Functional analysis
Resistance genes, plasmids, virulence factors, insertion sequence elements (IS), and phages were annotated using BLASTn e BLASTX against several specific databases, including CARD, ARDB, PlasmidFinder, ResFinder, VirulenceFinder, ICEberg, VFDB, ISFinder, phySPY, PHASTER, and PLSDB [7,22–26]. Outputs have been parsed to discard all the hits with a BIT score <300, coverage <65%, and similarity <70%. The remaining hits have been ordered by position, and in case of overlapping matches, the ones with the highest BIT score have been selected.

Metagenomics
Sequencing data pre-processing of the donor’s blood sample was performed by FASTP 0.20.1. Human reads were mapped against the human reference genome hg19 using bowtie2 and removed by samtools tools [27]. The host-filtered microbial reads were taxonomically profiled using MetaPhlAn2 (version 2.7.5) [28,29].

Results
Patients’ characteristics
In September 2020, four paediatric onco-hematologic patients received PCs from the same irradiated sample, prepared by a single-donor (a healthy 59-year-old Italian female) plateletpheresis at Bambino Gesù Children’s Hospital (OPBG), Rome, Italy.

Three patients were males, with a median (interquartile range, IQR) age of 8 (7–9) years (Table 1). Three of them were affected by acute lymphocytic leukemia (ALL) (patients 1–3), while one was affected by metastatic neuroblastoma (patient 4). All were immunocompromised (Table 1).

Approximately 24 h after transfusion, body temperature, C-reactive protein, and procalcitonin levels have been altered in all four patients, reaching elevated values (>38°C, >5 mg/dL, and >60 ng/mL, respectively) in three out of four patients (patients 2–4). Accordingly, a clinically defined sepsis was diagnosed in patients 2, 3, and 4 (Table 1 and Figure 1), while patient 1 experienced an acute self-limited illness. An empiric antimicrobial therapy mainly composed of tigecycline and aminoglycosides (Table 1) was administered to each patient as soon as clinical manifestations were evident. In all patients, the vital signs returned to physiological ranges in a median (IQR) time of 4.5 (3.5–5.75) days.

Phylogenetic relatedness of L. garvieae isolates
L. garvieae was isolated from the blood culture of all three patients developing sepsis (patients 2–4) 24 h post PCs transfusion, confirming the bloodstream infection. The blood culture of patient 1 resulted negative, confirming the self-limited illness.

The antimicrobial susceptibility testing defined resistance to the lincomamide clindamycin and penicillin G for all isolates (minimal inhibitor concentration [MIC] values: >256 and ≥0.5, respectively).

WGS experiment returned a total of 4.15 GB of Illumina 2 × 150 bp data by the sequencing of three L. garvieae isolates. De novo genome reconstruction retrieved an average genome size of 2.06 million base pairs with a GC content of 37.9%. The three L. garvieae sequences are available in the European Nucleotide Archive (ENA) under the following BioSample accession numbers: SAMEA12289652, SAMEA12289653, and SAMEA12289654.

Looking at the genome content, the three strains were characterized by 440 common SNPs, 44 of them were never described before (Table S2). Of these 43 SNPs, 19 were non-synonymous, and thus, involved in amino acid modifications in different L. garvieae enzymes.

The ML tree (based on the core SNP genomes of the three pediatric L. garvieae isolates plus 24 publicly available L. garvieae and L. lactis sequences, Table S1) showed that the three isolates shared a 99.9% identity, and clustered together within a bootstrap of 100% in the subtree of human and cow L. garvieae strains (Figure 2(A)). The common origin of the three L. garvieae strains and their strong relatedness with human and cow strains were further confirmed by the Bayesian inference as suggested by the topology of the tree and the clusters’ posterior probabilities equal to 1.00 (Figure 2(B)).

Of note, the isolates described in this work exhibit a genetic relatedness with L. garvieae isolates from meat and dairy products made with raw milk, previously
pointed as the potential sources of human \textit{L. garvieae} infection [5].

**Virulence factors and resistance genes characterizing the \textit{L. garvieae} isolates**

The isolates hold several virulence factors, such as hemolysin III family proteins, adhesin gene clusters, siderophores, and sortases (Table 2), suggesting a role of these strains in contributing to the pathogenic manifestations and the reported illnesses of the three pediatric patients. Antimicrobial resistance profile is promoted in the three isolates by chromosomal \textit{lsa(D)} gene, recently detected in \textit{L. garvieae} fish pathogenic strain and described as a novel factor for resistance to lincosamide, pleuromutilins, and streptogramins [30]. It acts as an ATP-dependent active transport and does not require additional cofactors to confer full resistance to upper mentioned antibiotics. The multidrug transporter \textit{Mdt(A)} gene, originally described as a plasmid-dependent antimicrobial resistance factor in \textit{E. coli} and \textit{L. lactis} [31], and recently found in the genomic content of \textit{L. garvieae} [32], was also shared in all the three isolates. Different from \textit{E. coli} and \textit{L. lactis}, this enzyme does not confer decreased susceptibility to erythromycin or tetracycline in \textit{L. garvieae} species, probably due to two amino acid mutations in the C-motifs of \textit{Mdt(A)}, known to suppress the resistance phenotype of Tet(K) and Tet (L) [33,34].

The intermediate penicillin resistance found by the antimicrobial susceptibility test was probably promoted to a group of chromosomal penicillin binding proteins (PBP1a, PBP1b, PBPs, PBP2a, PBP2b, Table 2) normally involved in wall cell biosynthesis [35], and thus, capable of reacting with β-lactams. The intermediate, instead of being fully resistant to penicillin, can be explained by the absence of an auxiliary set of protein and regulatory elements encoded by a cassette chromosome called \textit{mec} found in most of the penicillin-resistant \textit{Staphylococci}, but absent in other non-pathogenic species [36].

Of note, the genome content analysis also revealed a vanZ-like domain, known to be present in the genomes of clinically relevant bacteria, such as \textit{Bacillus}, \textit{Streptococcus}, \textit{Enterococcus}, and \textit{Clostridium}, and decreases their sensitivity to some lipoglycopeptide antibiotics, but not vancomycin [37,38].

### Table 1. Pediatric patients’ characteristics.

| Patient | Sex   | Age, years | Nationality | Disease | PCs transfusion (day/month/year; hour: minute) | At PCs transfusion: | Symptoms after positive blood culture: | Empiric antimicrobial therapy |
|---------|-------|------------|-------------|---------|---------------------------------------------|---------------------|----------------------------------------|---------------------------------|
| 1       | Female| 8          | Italian     | ALL     | 01/09/2020; 16:53                           | Yes                 | Fever, celsius 39.2                     | INN-Tigecycline; Amikacin; β-lactams |
| 2       | Male  | 9          | Italian     | ALL     | 02/09/2020; 14:04                           | Yes                 | Chills Yes                             | INN-Tigecycline; Amikacin; INN-ceftazidime/avibactam |
| 3       | Male  | 4          | Italian     | ALL     | 02/09/2020; 14:32                           | Yes                 | Heart rates, bpm 110                   | INN-Tigecycline; Amikacin; INN-ceftazidime/avibactam |
| 4       | Male  | 9          | Italian     | Metastatic neuroblastoma                    | 02/09/2020; 16:36   | Yes                                    | INN-Tigecycline; Amikacin; β-lactams; Quinolones; Metronidazole; |

ALL: Acute lymphocytic leukemia; PCs: Platelet concentrates; Hb: Hemoglobin; Ht: Hematocrit; MCV: Mean Corpuscular Volume; MCH: Mean Corpuscular Hemoglobin; MCHC: Mean Corpuscular Hemoglobin Concentration; RDW: Red Cell Distribution Width; HDW: Hemoglobin Distribution Width; CRP: C-reactive Protein; INN: International Non-proprietary Name. *Replaced by Gentamycin one day later.
Non-chromosomal genomic content in our isolates is represented by the plasmid pUC11C (Table 2) [39], known to encode two class C sortases, which are commonly involved in pilus biosynthesis [40,41]. The three L. garvieae genomes also contained the PLg-TB25 temperate prophage (coverage: 71.19%), recently described as a non-virulent prophage from a dairy strain of L. garvieae [42], thus confirming the genetic relatedness of the three L. garvieae strains highlighted by the phylogenetic tree. This prophage carries with it several enzymes, such as resolvase and helicase, promoting exogenous DNA integration into the bacterial chromosome, and thus, paying an active role in acquiring and fixing genetic elements going under positive selection.

**Donor’s platelet concentrates characterization**

To confirm the source of L. garvieae infection, the microbial reads obtained by the donor’s PCs were profiled – thanks to MetaPhlAn2 (v.2.7.5). After removing low-quality and host genome reads, we obtained 5,459,177 reads with a quality score >30. The 4.98% (number of reads: 271,810) were properly classified as bacteria (n = 64,145), viruses (n = 200,356), or eukarya (n = 7,309), while 95.2% remained unclassified. Among the bacterial reads, 29.9% (number of reads: 19,154) belonged to Firmicutes, and 5.16% (number of reads: 3,310) defined Streptococcaceae. The remaining bacterial reads mainly belonged to Proteobacteria (number of reads: 38,127).
| Gene product                          | Accession number and locus | Identity (%) | Length (nt) | E-value* | BitScore** |
|--------------------------------------|---------------------------|--------------|-------------|----------|------------|
| Chromosomal content                  |                           |              |             |          |            |
| Hemolysin III related protein        | NC_017490 region:331642-352295 | 98.61        | 217         | <1E-100  | 424        |
| U32 family peptidase [Collagenase]   | NC_017490 region:1699976-1701265 | 99.53        | 429         | <1E-100  | 887        |
| MucB domain-containing protein LCGL 1005 | AFCC01000004 region:18820-20355 | 97.65        | 680         | <1E-100  | 1321       |
| WxL surface protein                  | AFCD01000002 region:17404-18222 | 89.35        | 629         | <1E-100  | 1019       |
| LpxTG surface protein (Cna B-domain) (orf 25) | HM852551 WP_019291416.1 | 99.81        | 2115        | <1E-100  | 4031       |
| Sortase A                            | NC_017490 region:1300351-1306046 | 96.29        | 243         | <1E-100  | 443        |
| Adhesins gene cluster - WxL surface protein | NC_017490 region:878477-872476 | 96.55        | 203         | <1E-100  | 389        |
| Adhesins gene cluster - Cell surface putative DUF916 and DUF3324 domain containing protein | DUF916 and DUF3324 | 93.17        | 337         | <1E-100  | 629        |
| Adhesins gene cluster - Putative surface protein (COG4713 domain) | HM852557 AFCC01000002 region:19917-21412 | 98.95        | 190         | <1E-100  | 382        |
| Pili specific sortase SpaH/EbpB family LPxTG-anchored major pilin | WP 100222480.1 in AFCD01000005 region: 28828-35485 | 92.64        | 258         | <1E-100  | 460        |
| ABC transporter ATP-binding protein LCGL 1621 | NC_017490 region:159432-1596948 | 92.62        | 339         | <1E-100  | 565        |
| ABC transporter substrate-binding protein LCGL 1622 | NC_017490 region:544393-548003 | 92.01        | 313         | <1E-100  | 588        |
| Iron ABC transporter permease LCGL 0527 | NC_017490 region:449047-449570 | 93.98        | 316         | <1E-100  | 429        |
| Iron ABC transporter permease LCGL 0528 | NC_017490 region:155884-155999 | 94.90        | 294         | <1E-100  | 516        |
| Iron-ABC transporter permease LCGL 0529 | NC_017490 region:289210-289818 | 98.40        | 313         | <1E-100  | 605        |
| Iron-hydroxamate ABC transporter LCGL 0530 | NC_017490 region:1300531-1306046 | 98.40        | 313         | <1E-100  | 605        |
| Enolase [phosphopyruvate hydratase]   | NC_017490 region:106587-109166 | 97.11        | 316         | <1E-100  | 429        |
| Glyceraldehyde-3-phosphate dehydrogenase LCGL 19790 | NC_017490 region:1928003-1929013 | 100.00       | 336         | <1E-100  | 671        |
| Superoxide dismutase LCGL 0285 | NC_017490 region:289210-289818 | 99.01        | 202         | <1E-100  | 385        |
| FAD-dependent oxidoreductase LCGL 0664 | NC_017490 region:76821-769758 | 99.10        | 445         | <1E-100  | 907        |
| Glycoside hydrolase 1 protein        | NC_CP065637 region:716031-717467 | 99.88        | 478         | <1E-100  | 487        |

Putative virulence factors were annotated using BLASTn e BLASTX. Identity was estimated against the reference genome for each gene product. Only hits with a bit score >300, coverage >65%, and identity >80% were considered. *LSa(D) is a specific resistance factor providing tolerance to lincosamides, while mtd(A) is a drug antibiotic responsible for aspecfic resistance to tetracyclines and macrolides; both of them belong to chromosomal genetic content. 

**Penicillin binding protein (PBP) is a chromosomal genetic content that are normally involved in cell wall biosynthesis; their full resistance mechanism is achieved by the actions of auxiliary genes mainly found in Staphylococcus pathogenic strains (holding chromosomal mec cassette); they provide only partial resistance in other bacterial genera. The insertion sequences 6 family have been characterized as important factors determining bacterial genome shaping. In fact, they are responsible for exogenous genetic content integration via IS6 family transposase action (found to be part of plasmid PUC11C). E-value index returns the probability to get a match by chance. **BitScore is the match reliability index.

All Streptococcaceae reads (ENA accession number: ERS9886497) shared a >97% homology with L. garvieae isolates, thus confirming the presence of L. garvieae in the PCs transfusion.

**Discussion**

Here, we describe three cases of sepsis-related to the drug-resistant L. garvieae transmission in three onco-hematologic pediatric patients caused by a platelet transfusion obtained by the same healthy adult donor. The fourth patient, who had received a transfusion first, developed a self-limited illness, accompanied by a blood culture that remained negative. To our knowledge, this is the first report that described a clinically defined sepsis in a pediatric setting caused by this emerging human pathogen during a blood transfusion procedure.

The cases of L. garvieae infection in humans described so far are characterized by a favourable clinical course and regard manifestations such as endocarditis, septicemia, urinary tract infection, peritonitis, and liver abscess [3,9,11]. L. garvieae was also the cause of bacterial contamination of PCs [43,44],
and thus, represents a serious problem in transfusions, as demonstrated by the first case of sepsis caused by this transfusion-transmitted pathogen [43].

In this regard, the safety of the blood supply, including bacterial contamination of platelet products and transfusion-transmission risks associated with emerging pathogens [45,46] continues to represent a challenge for clinical blood centers. In the pediatric setting, sepsis without source account for 3.4%–13.6% of cases seen in emergency departments [47], and all are characterized by a challenging diagnosis. Today, there are certain technologies that mitigate this risk either as commercially available products or as investigational protocols. These include pathogen-reduction technology (PRT) for apheresis platelets and plasma, rapid tests for bacterial detection in PCs, and investigational screening assays for emerging pathogens. The implementation of these technologies has enhanced the safety of the blood supply in the last years [48], even if full prevention of transfusion-related bacterial infection cannot be completely achieved. In 2019, according to the latest FDA report, approximately 1.9 million apheresis platelets were transfused, and one death due to bacterial contamination occurred [49].

Here, the three cases of sepsis in pediatric recipients developed 24 h after PCs transfusion. All three patients (patients 2–4) were characterized by a peak fever and a significant C-reactive protein and procalcitonin increase (Figure 1). Consistent with clinical manifestations, the blood culture revealed L. garvieae infection in all three patients. The genome content analysis of L. garvieae isolates from the three blood cultures suggested their clonal origin and a well-defined homology with L. garvieae derived from meat and dairy products [7]. The source and the transmission chain were revealed by the metagenomics analysis that confirmed the presence of L. garvieae traces in PCs.

Even if all patients had a favourable clinical course after a fully active antimicrobial therapy composed mostly of tigecycline and aminoglycosides (Amikacin), the illnesses reported in three out four patients after transfusion were typical of a pathogenic microorganism invasion.

In view of this and other reports that described sepsis associated with L. garvieae infection [11,13,50,51], the knowledge of virulence factors and resistance mechanisms associated with the Lactococcus genera and L. garvieae species is detrimental.

Here, we implemented the knowledge of the virulent L. garvieae circulating strains by providing a capillary description of chromosomal and extrachromosomal content of these bacteria, focusing the attention to all factors that might confer a selective advantage in host invasion [7]. Indeed, the three strains hold the virulence factors necessary to survive and feed in iron rich-environment, like human blood [52]. Collected strains also shared several groups of adhesins, haemolysin, fibronectin-binding proteins, and penicillin acylase that actively promote bacterial colonization of mucosal tissues [53,54], and thus, increase the chance of bacteria being present in blood transfusable components [52,54]. Some adhesins such as MucBP domain-containing protein LCGL 1005 also confer to the bacteria the ability to form biofilm and to escape immune surveillance systems [55]. Furthermore, the presence of WxL domain-containing proteins (already characterized in Enterococcus faecium) increases the L. garvieae ability to overcome the osmotic stress and to aggregate in a complex population [56].

We also identified chromosomal contents conferring drug resistance to lincosamides (IsaD gene) and penicillins (penicillin binding proteins) (Table 2), and to some lipoglycopeptide antibiotics [31,35,37,38,57], thus providing the genetic basis of the antimicrobial susceptibility testing results, that defined resistance to clindamycin and intermediate resistance to penicillin G. These results also confirmed the drug-resistant genetic backbone of the L. garvieae isolated in the three pediatric patients.

Even if the three cases described here support the circulation of drug-resistant L. garvieae strains in humans and its potential role as a human pathogen, more insights and evidence are needed to better define L. garvieae pathogenicity and related outcomes, and thus, to guide its significance in clinical practice. Most efforts in WGS are also needed to better characterize L. garvieae circulation in pediatric and adult settings.

Regarding transfusion-transmitted infection risk, although four patients had received the same platelet concentrate bag, only three developed sepsis. A discriminating factor that can justify the different grades of illness is the time of transfusion administration.

According to our hospital procedure [58], PCs are generally stored on a platelet shaker at room temperature (22 ± 2°C) before being transfused. Due to the ability of L. garvieae to grow at temperatures as low as 10°C in red blood cells [59], the platelets storage at room temperature could have provided suitable conditions for bacterial proliferation [48].

In line with this evidence, we observed that the first patient who received the transfusion was the only one who did not develop sepsis. On the contrary, the patient who last received the platelet concentrate bag was the first to have a positive blood culture. Interestingly, this patient was also characterized by a higher level of procalcitonin than the others (Figure 1), providing further evidence that in the case of bacterial contamination, the longer is the time of platelets storage, the higher is the risk to transmit high bacterial load [60]. It is worth knowing that the irradiation
performed according to internal hospital procedures is used to prevent the Graft *versus* Host Disease (GvHD), but is not able to prevent bacterial growth as stated in the National Guidelines [58].

In conclusion, despite advances in donor screening and infectious disease testing, the risk of transfusion-transmitted infections continues to be of particular concern, as defined by the three episodes of sepsis due to a transfusion-associated transmission of drug-resistant *L. garvieae* in pediatric hospitalized oncologic patients. This highlights the importance to implement the screening of platelet components with new human-defined pathogens to ensure the safety of blood supply, and more broadly, for surveillance of emerging pathogens.

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**Disclosure statement**

No potential conflict of interest was reported by the author (s).

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