Hospital outbreak caused by linezolid resistant \textit{Enterococcus faecium} in Upper Austria

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Abstract

**Background:** Enterococcus faecium is part of the human gastrointestinal flora but may act as opportunistic pathogen. Environmental persistence, high colonization capability and diverse intrinsic and acquired resistance mechanisms make it especially successful in nosocomial high-risk settings. In March 2014, an outbreak of Linezolid resistant Enterococcus faecium (LREfm) was observed at the hematooncology department of a tertiary care center in Upper Austria.

**Methods:** We report on the outbreak investigation together with the whole genome sequencing (WGS)-based typing results including also non-outbreak LREfm and susceptible isolates.

**Results:** The 54 investigated isolates could be divided in six clusters based on cgMLST. Cluster one comprised LREfm isolates of genotype ST117 and CT24, which was identified as the causative clone of the outbreak. In addition, the detection of four other clusters comprising isolates originating from hematooncology patients but also at other hospitals, pointed to LREfm transmission between local healthcare facilities. LREfm patients ($n=36$) were typically at risk for acquisition of nosocomial pathogens because of immunosuppression, frequent hospitalization and antibiotic therapies. Seven of these 36 patients developed LREfm infection but were successfully treated. After termination of the initial outbreak, sporadic cases occurred despite a bundle of applied outbreak control interventions.

**Conclusions:** WGS proved to be an effective tool to differentiate several LREfm clusters in an outbreak. Active screening for LREfm is important in a high-risk setting such as hematooncology, where multiple introductions are possible and occur despite intensified infection control measures.

**Keywords:** Enterococcus faecium, Linezolid, Austria, Whole genome sequencing

Background

Enterococci are gram-positive bacteria found in the environment and as part of the human gastrointestinal flora [1]. They can act as opportunistic pathogens causing a broad range of diseases such as blood stream or wound-associated infections [2]. Hospital-adapted clones such as clonal complex (CC)17 \textit{Enterococcus faecium} show persistence in the environment and high colonization capability [3]. \textit{E. faecium} has diverse intrinsic resistance mechanisms to antibiotics and is able to progressively acquire antimicrobial resistances such as to ampicillin and vancomycin (VAN), thus limiting the therapeutic options [4]. One therapy of last resort against vancomycin-resistant Enterococci (VRE) is the oxazolidinone linezolid (LZD), which inhibits protein synthesis by binding to the 50S 23S rRNA [1]. Resistance to LZD has already been reported in \textit{E. faecium} with the most common resistance mechanisms referring to mutations in the V domain of the 23S rRNA [5, 6]. Mutations in the sequence of genes encoding the riboproteins L3, L4 and L22 account for the second most common mechanisms. Third, recently described plasmid-mediated resistances due to \textit{cfr} [7], \textit{optrA} [8] and \textit{poxA} [9] and finally, yet unknown LZD resistance mechanisms are known to exist [10]. Risk factors associated with LZD resistance

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include transplants and surgery, immunosuppression and previous or ongoing treatment with LZD [11, 12]. Usually, LZD resistant E. faecium (LREfm) strains emerge in patients 22–125 days after treatment [13]. LREfm outbreaks tend to be mostly colonizations, although clinical outbreaks with invasive LREfm infections have been reported [14].

**Methods**

In March 2014, routine surveillance cultures of stool and urine from four patients hospitalized at the department of internal medicine 1 (DIM1) of an Austrian tertiary care center tested positive for LREfm. An outbreak investigation was then initiated to identify the source of the outbreak, characterize the outbreak strain by whole genome sequencing (WGS) and antimicrobial susceptibility testing and apply control measures.

**Hospital and ward description**

The DIM1 is one of three departments of internal medicine at a tertiary care center (hospital 4) in Linz, Upper Austria. It consists of a general oncology ward (16 rooms for 33 patients), a leukemia and autologous stem cell transplantation ward (8 rooms for 12 patients), a stem cell transplantation unit (5 rooms for 5 patients) and an outpatient clinic. The bed occupancy rate ranges between 90 and 100%.

**Outbreak description**

Between March and May 2014, we identified ten LREfm colonized patients and one patient with LREfm bloodstream infection who were all hospitalized at the DIM1. The outbreak was contained in June 2014 through implementation of control measures. Active case finding detected twelve additional patients at the DIM1 and four at other units of hospital 4 till the end of the year 2017 (total number of cases: 27). Thirteen of these 16 patients were categorized as colonized, one patient treated at DIM1 had LREfm bloodstream infection and two patients treated in other departments had a urinary tract infection and a surgical site infection, respectively. A case was defined as any patient with culture-confirmed LREfm identified at hospital 4 from the beginning March 2014 onwards. Cases were included in this study until the end of 2017.

**Patients**

In total, 36 patients were included in the study. Twenty-seven of those were the cases from hospital 4 as described above (24 were patients of the DIM1 and three were patients from other wards), who were designated as patient collective A. In order to gain insight into the occurrence of the outbreak strain in the local and Austrian hospital community, an additional nine patients belonging to six other hospitals (hospitals 1 to 3 and 5 to 7) in the provinces of Carinthia, Upper Austria and Vienna were included in the study and denominated patient collective B. Their isolates had been collected between 2012 and 2018. Demographic and epidemiological data such as patient outcome (death/survival), immune status, routine screening or clinical sampling, hospital contact in the previous year and other parameters were collected, anonymized and then analyzed using Microsoft Excel 2016 and SPSS statistical software version 22.0 (Chicago, Illinois). Data on the exposure to LZD in the 28 days prior to isolation of LREfm and current antibiotic treatment (yes/no) was retrieved from patient records at the time of the first LREfm isolation. If available, duration of LREfm carriage was estimated from surveillance culture data. LZD consumption data of the DIM1 were extracted from the hospital pharmacy database and analyzed by AVS.webKess software [15]. We investigated at patient level (patient collectsives A and B, n = 36) the acquisition of the outbreak strain and possible associations with their demographics and epidemiological characteristics using STATA 13 software.

**Isolates**

Isolates (n = 54) were retrieved from cryobanks (Mast, Reinfeld, Germany) and cultured on blood agar (Oxoid, Wesel, Germany). Maldi-TOF analysis (Bruker Daltonics, Bremen, Germany) was used for species confirmation. LZD susceptibility testing was done using disk diffusion and, for the 45 LREfm isolates, gradient testing (Biomerieux, Marcy L’Étoile, France) according to EUCAST criteria and breakpoints (resistant: > 4 mg/L or < 19 mm, respectively). Additionally, Tedizolid (TDZ) gradient testing was performed for all LREfm (Liofilchem, Roseto degli Abruzzi, Italy) according to the manufacturer’s instructions. VAN susceptibility data were retrieved from the routine antibiogram. Finally, all LREfm were investigated for the presence of optrA and cfr using previously published primer sets [16].

For comparison of WGS-based typing data, we included nine additional LZD susceptible E. faecium (LSEfm) isolates (eight from blood culture and one from stool) recovered from eight patients staying at hospitals 4 and 6 between May 2014 and June 2017 (patient collective C). Two of those were collected from patients having also an LREfm isolate. The others were chosen because they were invasive isolates from affected wards.

**DNA extraction, WGS and typing**

High-molecular-weight DNA from the 54 bacterial overnight cultures was isolated using a MagAttract HMW DNA kit (Qiagen, Hilden, Germany). DNA was quantified using DropSense 16 (Trinean NV/SA, Gentbrugg, Belgium). Library for WGS was prepared with a NexteraXT kit (Illumina, Inc., San Diego, CA, USA) according to manufacturer’s instructions and a 300-bp paired-end sequencing run was performed on an Illumina MiSeq instrument using the MiSeq
V3 reagent kit (Illumina Inc., San Diego, CA, USA) for the 54 E. faecium isolates. Raw reads were de novo assembled into draft genomes using SPAdes version 3.11.1 [17]. Contigs were then filtered for a minimum coverage of 5 and minimum length of 200 base pairs. SeqSphere+ software (Ridom GmbH, Münster, Germany) was used for strain typing using a public core genome multilocus sequence typing (cgMLST) scheme [18]. Minimum spanning trees (MST) were generated to illustrate the number of allelic differences between isolates and visualize clusters. The allelic cluster threshold was set to ≤20 allelic differences as previously proposed [18]. Sequence types (STs) from the classical MLST [19] were in silico extracted from WGS data using SeqSphere+. Likewise, the three genes encoding the riboproteins L3, L4 and L22 and the 23S rRNA genes were extracted and screened for point mutations associated to LZD resistance by comparing their sequences with those of the E. faecium DO reference strain Antimicrobial resistance genes including those conferring LZD resistance (optrA, postA and cfr and its variants) were identified via the Comprehensive Antibiotic Resistance Database (CARD) [20]. In addition, both the point mutations and the genes conferring LZD resistance including G2576U located in the 23S rRNA were double-checked using LRE finder [21]. The presence of specific virulence genes (VGs) for E. faecium among the isolates was investigated using the Virulence Factors Database and Virulence Finder 2.0 [22, 23]. Plasmid Finder was used for plasmid identification among the sequenced isolates [24]. Lastly, we assessed at an isolate level (n = 54) possible associations between the resistance phenotype (LREfm/LSEfm) of the isolates and the presence of VGs, resistance genes and the point mutation 23S rRNA G2576U.

**Results**

**Characteristics of all study patients and epidemiological data**

In the study period a total of LREfm 36 patients were detected (patient collectives A and B). Their demographic and epidemiological characteristics are summarized in Table 1. Figure 1a shows the epidemiological curve of the outbreak by collection date of the first LREfm isolate. Patients with more than one isolate are represented once only in the epicurve. The location of the hospitals is shown in Fig. 1b.

**Patient movement and clinical characteristics**

When patient movement of cases (collective A) was traced, possible direct transmission could be shown for 10 out of 11 cases during the initial outbreak event (March 2014 to May 2014). Five of the eleven cases shared a room with confirmed LREfm cases and another five were treated in wards at the same time as confirmed cases (Fig. 2).

Of the 16 additional cases detected between September 2014 and November 2017, six shared at least the DIM1

| Table 1 Clinical and epidemiological data obtained for the 36 patients of collective A and B |
|---------------------------------|-----------------|-----------------|
| **Characteristics**             | **Age, years, median (range)** | **Sex, n (%)** |
|                                 | 57 (22–91)      | Male            |
|                                 |                 | 23 63.9         |
|                                 |                 | Female          |
|                                 |                 | 13 36.1         |
| Patient status, n (%)           | Inpatient       | 35 97.2         |
|                                 | Outpatient      | 1 2.8           |
| Transfer from other Hospital/LTCF, n (%) | Yes | 8 22.2 |
|                                 | No              | 25 69.4         |
|                                 | Unknown         | 3 8.3           |
| Duration of LZD therapy, days, median (range) | 9 (3–35) |
| LREfm status, n (%)             | Infection       | 4 11.1          |
|                                 | Colonization    | 30 83.3         |
|                                 | Unknown         | 2 5.6           |
| Duration of colonization, days, median (range) | 22 (3–245) |
| Current antibiotic therapy, n (%) | Yes | 36 100.0 |
|                                 | No              | 0 0.0           |
| Hospital contact in previous year, n (%) | Yes | 31 86.1 |
|                                 | No              | 2 5.6           |
|                                 | Unknown         | 3 8.3           |
| Underlying disease, n (%)       | Hematology-Oncology | 25 69.4 |
|                                 | allogeneic SCT  | 17              |
|                                 | autologous SCT | 3               |
|                                 | Trauma/orthopedic | 4 11.1 |
|                                 | Abdominal surgery/Pancreatitis | 4 11.1 |
|                                 | Other           | 3 8.3           |
| Immunocompromised, n (%)        | Yes            | 31 86.1         |
|                                 | No              | 4 11.1          |
|                                 | Unknown         | 1 2.8           |
| Died during follow-up (until end of 2017), n (%) | Yes | 15 41.7 |
|                                 | No              | 14 38.9         |
|                                 | Unknown         | 7 19.4          |
with known LREfm carriers during their hospitalization. For ten cases, no direct epidemiological link could be established except for possible indirect transmission on previously affected wards.

Most patients (collective A + B) had multiple LREfm isolates, typically from stool and urine, but eventually lost those enterococci again. To estimate duration of colonization, time from first LREfm isolate until clearance was available for 21 patients and ranged from three to 245 days (median 22 days). For only two patients the same LREfm clone could be cultured again after more than 1 year.

Five patients staying at hospital 4, one at hospital 6 and one at hospital 1 had an LREfm infection (total n = 7), the rest was considered colonized. All seven infected patients were treated with VAN and none of them died. Fifteen (41.6%) patients died during follow-up but only two died during their stay at hospital 4 at the time of LREfm detection of unrelated causes. All patients (100%) received antibiotics at the time of LREfm detection. Ten

### Table 1 Clinical and epidemiological data obtained for the 36 patients of collective A and B (Continued)

| Characteristics                  | Source material first LREfm isolate, n (%) | Source material secondary isolates, n (%) | Screening sample, n (%) |
|----------------------------------|-------------------------------------------|------------------------------------------|-------------------------|
|                                  | Stool 18 (50.0)                           | Blood culture 2                          | Yes 24 (66.6)           |
|                                  | Urine 9 (25.0)                            | Urine 1                                  | No 11 (30.5)            |
|                                  | Swab (wound, throat, eye) 7 (19.4)        |                                          |                         |
|                                  | Blood culture 1 (2.8)                     |                                          |                         |
|                                  | Catheter tip 1 (2.8)                      |                                          |                         |

#### Fig. 1

**A.** Epidemiological curve representing the 36 patients with a LZD resistant isolate within the period 2012–2018. **B.** Map of Austria with locations of the seven hospitals that provided LREfm isolates.
(27.8%) patients had received LZD in the 28 days prior to the recovery of LR_Efm, whereas 25 had not been exposed. Eight patients (22.2%), all from hospital 4, had been transferred from other hospitals, two of them from hospital 6. Thirty-one (81.1%) patients had been hospitalized in the previous year and the same number (81.1%) were immunocompromised. The majority of patients (n = 24, 66.6%) had been tested for LR_Efm as part of screening.

**Antimicrobial susceptibility testing**

The LR_Efm isolates were all LZD resistant by disk diffusion testing and had a median MIC of 32 mg/L (range 4 to > 256). The two isolates with an MIC of 4 mg/L were included because of their LZD zone diameters of 17 and 18 mm, respectively. TZD MICs ranged from 1 to 32 mg/l (median 2 mg/L) and were all lower than the respective LZD MICs. Three isolates showed MICs of > 256 mg/l for VAN.

**WGS-based typing**

WGS-based typing assigned the 54 isolates to seven sequence types (STs) (Table 2), with ST117 (n = 30; 55.5%) and ST80 (n = 18; 33.3%) being the most frequent. Within the LR_Efm group (n = 45), ST117 was the predominant ST (n = 28, 62%) while for the LS_Efm ST80 was the most frequent ST (n = 5, 56%). The vast majority of ST117 isolates belonged to cluster type (CT) 24 (n = 25, 83%). All the other STs were found once or twice. Based on the cgMLST data of the 54 isolates, the MST revealed six clusters (Fig. 3). We only identified one cluster (cluster 1) with 25 LR_Efm isolates (46.3%, from 20 patients overall) as part of the outbreak at the DIM1 in hospital 4. An LS_Efm isolate from a case also clustered in cluster 1. All isolates within the cluster were ST117 and all but one presented CT24. Isolates of cluster 1 differed by 0 to 10 alleles.

WGS-based typing revealed that 10 out of 11 cases (see also Fig. 2, all patients except patient 30 / Ef-13) detected at the beginning of the outbreak were part of it and therefore their isolates (n = 15) belonged to cluster 1, confirming the transmission suspected by patient movement data. After May 2014, nine cases were detected and these also carried the outbreak clone (n = 10 isolates, grouping in cluster 1). Last, seven additional cases from hospital 4 detected after May 2014 were not part of the outbreak, grouping five LR_Efm isolates in clusters 2 (n = 2) and 3 (n = 3). At a patient level (n = 36), we found significant associations between acquisition of the LR_Efm outbreak strain and screening sampling (Exact p < 0.005) as well as a fatal outcome during follow-up (Exact p < 0.005).

Clusters 2 to 6 comprised isolates from patients staying at different hospitals and from a wider period (2012–2018) (Fig. 3). Cluster 2 (ST80, CT1873) comprised two LR_Efm isolates from two patients of hospital 4, not treated at the DIM1, four LR_Efm isolates from four patients at hospitals 2 (n = 2), 3 (n = 1) and 6 (n = 1) and also one LS_Efm isolate from hospital 6. None of these patients was treated for an oncological disease, they were typically older (median age 69 years) and two had been transferred from long term care facilities.

Cluster 3 (ST80, CT16) consisted of five isolates from four patients treated at the DIM1, of which two were LS_Efm. Cluster 4 (ST80, CT315) grouped three isolates from three patients staying at hospitals 6 (n = 2) and 4 (n = 1), the latter being an LS_Efm from a patient who had not been treated at the DIM1. Cluster 5 (ST203, CT20) grouped two isolates from a patient staying at hospital 5 in Vienna. Cluster 6 (ST117, CT929) comprised two isolates obtained at two hospitals located in different regions. One of them originated from hospital.
| Case | Province | Hospital | Ward | Isolate | Isolation year | Source | Phenotype | LRE mechanism | ST | CT | Cluster number | Linezolid MIC (mg/L) | Tedizolid MIC (mg/L) |
|------|----------|----------|------|---------|----------------|--------|-----------|---------------|----|----|----------------|---------------------|---------------------|
| 1    | UA       | 2        | 2    | EF-03   | 2013 Wound swab | LREfm  | unknown   |               | 80 | 1873 | 2              | 32                  | 4                   |
| 2    | UA       | 4        | 5    | EF-11   | 2014 Stool    | LREfm  | G2576U, A2598G | 117 | 24  | 1               | > 256                | 32                  |
| 2    | UA       | 4        | 6    | EF-22   | 2014 Blood culture | LREfm  | unknown   |               | 117 | 24  | 1               | 32                  | 4                   |
| 3    | UA       | 6        | 13   | EF-44   | 2016 Wound swab | LREfm  | unknown   |               | 80 | 1873 | 2              | 64                  | 2                   |
| 4    | UA       | 4        | 10   | EF-36   | 2015 Wound swab | LREfm  | unknown   |               | 80 | 1873 | 2              | 128                 | 4                   |
| 6    | UA       | 4        | 5    | EF-17   | 2014 Urine    | LREfm  | G2576U   |               | 117 | 24  | 1               | 32                  | 8                   |
| 6    | UA       | 4        | 5    | EF-20   | 2014 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 64                  | 2                   |
| 7    | UA       | 4        | 5    | EF-15   | 2014 Urine    | LREfm  | unknown   |               | 117 | 24  | 1               | 16                  | 4                   |
| 8    | UA       | 4        | 11   | EF-43   | 2016 Throat   | LREfm  | G2576U   |               | 117 | 24  | 1               | 32                  | 2                   |
| 9    | UA       | 4        | 5    | EF-05   | 2014 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 16                  | 2                   |
| 10   | V        | 5        | N/A  | EF-24   | 2015 Stool    | LREfm  | unknown   |               | 203 | 20  | 5               | 16                  | 2                   |
| 11   | UA       | 4        | 5    | EF-40   | 2015 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 32                  | 2                   |
| 12   | UA       | 4        | 6    | EF-18   | 2014 Urine    | LREfm  | G2576U   |               | 117 | 24  | 1               | 32                  | 2                   |
| 13   | UA       | 4        | 5    | EF-37   | 2015 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 64                  | 2                   |
| 14   | UA       | 4        | 6    | 13639   | 2015 Blood culture | LSEfm  | unknown   |               | 117 | 24  | 1               | n.d.                | n.d.                |
| 14   | UA       | 4        | 6    | EF-31   | 2015 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 32                  | 2                   |
| 15   | UA       | 4        | 9    | EF-30   | 2014 Urine    | LREfm  | unknown   |               | 80 | 1873 | 2              | 64                  | 4                   |
| 16   | UA       | 4        | 5    | EF-28   | 2014 Urine    | LREfm  | G2576U   |               | 117 | 24  | 1               | 64                  | 8                   |
| 17   | UA       | 4        | 6    | EF-35   | 2015 Stool    | LREfm  | unknown   |               | 80 | 16  | 3               | 64                  | 4                   |
| 19   | UA       | 4        | 7    | EF-26   | 2015 Urine    | LREfm  | G2576U   |               | 117 | 1875 | not clustered | 64                  | 8                   |
| 20   | UA       | 4        | 6    | EF-23   | 2014 Stool    | LREfm  | G2576U, A2598G | 117 | 1872 | 1              | 32                  | 4                   |
| 20   | UA       | 4        | 6    | EF-27   | 2014 Stool    | LREfm  | G2576U   |               | 117 | 24  | 1               | 32                  | 16                  |
| 21   | UA       | 6        | 12   | EF-42   | 2016 Catheter | LREfm  | G2576U, A2598G | 80 | 315 | 4              | 64                  | 4                   |
| 22   | UA       | 2        | 1    | EF-02   | 2013 Wound swab | LREfm  | G2576U   |               | 80 | 1873 | 2              | 128                 | 16                  |
| 23   | UA       | 4        | 5    | EF-39   | 2015 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 4                   | 1                   |
| 24   | UA       | 4        | 6    | EF-29   | 2014 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 8                   | 2                   |
| 25   | UA       | 4        | 5    | EF-48   | 2017 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 16                  | 1                   |
| 26   | UA       | 4        | 5    | EF-46   | 2017 Stool    | LREfm  | unknown   |               | 117 | 24  | 1               | 32                  | 1                   |
| 27   | UA       | 4        | 5    | EF-19   | 2014 Urine    | LREfm  | unknown   |               | 117 | 24  | 1               | 32                  | 2                   |
| 27   | UA       | 4        | 5    | EF-21   | 2014 Stool    | LREfm  | unknown   |               | 80 | 1879 | not clustered | not clustered      | 4                   |
| 28   | UA       | 4        | 5    | EF-47   | 2017 Stool    | LREfm  | unknown   |               | 80 | 1876 | not clustered | 32                  | 2                   |
| 29   | UA       | 4        | 6    | 14921   | 2016 Blood culture | LSEfm  | unknown   |               | 80 | 16  | 3               | n.d.                | n.d.                |
| 29   | UA       | 4        | 5    | EF-32   | 2015 Stool    | LREfm  | unknown   |               | 80 | 16  | 3               | 64                  | 2                   |
7 in Carinthia and the other one was an LSEfm stool isolate from a DIM1 patient at hospital 4, from whom also three LREfm isolates were included in the study all clustering in cluster 3. The timespan between obtainment of cluster 6 isolates was 2 years and there was no epidemiological data to explain their relatedness.

Last, a number of isolates did not cluster with any other isolate: five LREfm from five patients at hospital 4, three LS
Efm from two patients at hospital 4 and one at hospital 6 and one isolate from one patient at hospital 1.

Twenty-four resistance genes were detected among the 54 isolates (Additional file 1: Table S1). The most common ones were AAC (6’)-li (n = 52; 96.3%), efmA (n = 52; 96.3%) and ermB (n = 48; 88.9%). Moreover, the multidrug efflux pump efmA was detected among all LREfm isolates within cluster 1. Significantly higher proportions were found for dfrF (Fisher’s exact P = 0.0020) and sat4 (Fisher’s exact P = 0.0499) within the LREfm (n = 45).

PCRs targeting cfr and optrA genes were negative for all LREfm. These genes, as well as poxtA, were also absent when blasting WGS data against the CARD database, the LRE tool from CGE server and Plasmid finder. Thirteen out of 36 (36.1%) cases presented at least one LREfm isolate carrying the point mutation G2576U at the 23S rRNA, as revealed when using CARD database meaning that 25.9% (n = 14) of the E. faecium isolates carried that point mutation (Table 2 and Additional file 1: Table S1). The number of 23S rRNA mutated copies varied between 2 and 6. In addition, three of those isolates carried a novel point mutation at A2598G of the

| Case | Province | Hospital | Ward | Isolate | Isolation year | Source | Phenotype | LREmechanism | ST | CT | Cluster number | Linezolid MIC (mg/L) | Tedizolid MIC (mg/L) |
|------|----------|----------|------|---------|----------------|--------|-----------|--------------|----|----|---------------|---------------------|---------------------|
| 29   | UA       | 4        | S    | EF-33   | 2015           | Blood culture | LREfm | unknown   | 80 | 16 | 3             | 64                  | 4                   |
| 29   | UA       | 4        | S    | EF-41   | 2016           | Stool     | LSEfm | unknown   | 117 | 929 | 6             | 2                   | 2                   |
| 30   | UA       | 4        | S    | EF-13   | 2014           | Stool     | LREfm | unknown   | 117 | 1878 | not clustered | 16                  | 2                   |
| 30   | UA       | 4        | S    | EF-14   | 2014           | Stool     | LREfm | G2576U    | 117 | 24 | 1             | 8                   | 2                   |
| 30   | UA       | 4        | S    | EF-34   | 2015           | Urine     | LREfm | unknown   | 117 | 24 | 1             | 64                  | 2                   |
| 31   | UA       | 4        | S    | EF-07   | 2014           | Urine     | LREfm | unknown   | 117 | 24 | 1             | 16                  | 2                   |
| 31   | UA       | 4        | S    | EF-08   | 2014           | Stool     | LREfm | unknown   | 117 | 24 | 1             | 16                  | 2                   |
| 32   | UA       | 4        | S    | EF-12   | 2014           | Stool     | LREfm | unknown   | 117 | 24 | 1             | 16                  | 2                   |
| 33   | UA       | 3        | S    | EF-04   | 2013           | Eye swab  | LREfm | G2576U    | 80  | 1873 | 2             | 32                  | 4                   |
| 34   | UA       | 6        | 16   | EF-38   | 2015           | Blood culture | LREfm | unknown   | 80  | 315 | 4             | 32                  | 2                   |
| 35   | UA       | 4        | S    | EF-09   | 2014           | Stool     | LREfm | G2576U    | 117 | 24 | 1             | 16                  | 2                   |
| 36   | C        | 1        | 1    | EF-01   | 2012           | Wound swab | LREfm | unknown   | 192 | 1877 | not clustered | 16                  | 2                   |
| 37   | UA       | 4        | S    | EF-45   | 2016           | Stool     | LREfm | unknown   | 78  | 1874 | not clustered | 32                  | 4                   |
| 38   | UA       | 6        | 15   | 14962   | 2016           | Blood culture | LSEfm | n.a.      | 1479 | not assigned | not clustered | n.d.                | n.d.                |
| 39   | UA       | 4        | 5    | 12981   | 2014           | Blood culture | LSEfm | n.a.      | 80  | 16 | 3             | n.d.                | n.d.                |
| 40   | C        | 7        | 14   | EF-51   | 2018           | Urine     | LREfm | G2576U    | 117 | 929 | 6             | 16                  | 2                   |
| 41   | UA       | 6        | 15   | 14522   | 2015           | Blood culture | LSEfm | n.a.      | 80  | 1873 | 2             | n.d.                | n.d.                |
| 42   | UA       | 4        | 4    | 15549   | 2016           | Blood culture | LSEfm | n.a.      | 1466 | not assigned | not clustered | n.d.                | n.d.                |
| 43   | UA       | 4        | 7    | 15378   | 2016           | Blood culture | LSEfm | n.a.      | 80  | 467 | 4             | n.d.                | n.d.                |
| 45   | UA       | 4        | 4    | 15925   | 2017           | Blood culture | LSEfm | n.a.      | 80  | 1873 | not clustered | n.d.                | n.d.                |

*Numbering allocated for each cluster in the MST
n.a. not applicable, n.d. not done
C Carinthia, CT Cluster type (cgMLST), MST minimum spanning tree, ST Sequence Type (classical MLST), UA Upper Austria, V Vienna
23S rRNA. For 31 LREfm isolates, the resistance mechanism was not found.

The three VAN resistant isolates (the two ST203 isolates from cluster 5 and a non-clustered ST117 isolate) carried the gene vanA.

Ten VGs were detected among the typed LREfm and LSEfm isolates (Additional file 1: Table S2). The most frequently found VGs were sgrA (n = 53, 98%), acm (n = 47, 87%) and ecbA (n = 33, 61.1%) and only ecbA was found to be significantly more frequent (Fisher’s exact P = 0.013) among LREfm isolates. We did not detect associations between ST and any of the VGs tested.

**Outbreak control measures**

After the detection of the outbreak, active surveillance was put in place for the DIM1 including antimicrobial susceptibility tests for all clinical and screening E. faecium isolates. In addition, strict contact precautions for known colonized patients, single-room patient care and increased frequency of cleaning and disinfection of patient-near-surfaces especially in bathrooms was initiated. Moreover, LZD consumption was reduced by preferential use of VAN for gram-positive coverage in empiric antimicrobial therapy of neutropenic patients (Fig. 4).

The outbreak was contained as of June 2014, however until the end of 2017 sixteen new cases were detected at hospital 4 at a rate of one per 2.6 months (Fig. 1). The fact that nearly half of these later cases were LREfm clones other than the outbreak clone could be revealed by WGS only retrospectively.

**Discussion**

In this study, we report the first LREfm outbreak in an Austrian hospital, which involved colonized as well as infected patients. WGS identified a hospital-adapted ST117, CT24 LREfm clone as the main causative agent of the outbreak in the affected hemato-oncology department. There was high clonality among this cluster, since all strains but one were ST117, CT24, and differed by
≤10 alleles. Interestingly, the outbreak clone did not spread to any other departments within hospital 4, meaning that the source was confined to DIM1. *E. faecium* ST117 is a widely disseminated clone that may present resistance to many drugs, including LZD [25]. In particular, ST117, CT24 has been previously identified in outbreaks caused by VAN-resistant *E. faecium* [26, 27] although up to our knowledge, this clone has been never associated to LR*Efm* outbreaks.

The addition of isolates from patient collectives B and C allowed us to gain more insight into the distribution of *E. faecium* between different healthcare facilities, especially for the region of Upper Austria. WGS also helped to elucidate that the outbreak at hospital 4 not just consisted of the cluster 1 clone (ST117, CT24), but also of two other clusters (2 and 3) and multiple unclustered isolates, suggesting several separate introductions of LR*Efm*.

Cluster 2 (ST80, CT1873) indicates a continuous spill-over between four healthcare facilities located in the same province even though patient movement could not be analyzed comprehensively. Cluster 3 (ST80, CT16) represented a separated cluster within the DIM1, in which a patient showed loss of LR*Efm* during a one year period. Similarly, in cluster 4 (ST80, CT315) we could observe transmission of *E. faecium* between hospitals 4 and 6, although the isolate detected at hospital 6 was LSE*fm*, which might indicate the loss of the resistance trait in this strain. Concerning cluster 6 (ST117, CT929), available epidemiological data do not allow to understand why two isolates from different time periods and geographical areas clustered together.

The clustering together of LR*Efm* and LSE*fm* in nearly all the clusters is an interesting finding in our study and, to our knowledge, it has not been described previously. However, outbreak reports usually only include resistant isolates [28]. An explanation for our finding of genetically closely related LR*Efm* and LSE*fm* isolates may be the different antibiotic selective pressure in diverse hospitals.

All LR*Efm* cases received concurrent antibiotic treatment, which may account for selection of enterococci in the enteric microbiome. Recent exposure to LZD was found only in a minority (n = 10; 27.8%) of our cases, however, highlighting the contribution of environmental and person-to-person spread as previously described [25]. This is supported by the fact that for the majority of DIM1 cases close physical contact could be confirmed. Hemato-oncological patients generally are at high risk for infections, but with profound immunosuppression, frequent hospitalization and repeated administration of antibiotics they are especially prone to acquisition of multidrug-resistant *E. faecium* [29]. The association of acquisition of the outbreak strain with screening and death during follow-up, however, is most likely confounded by the intrinsic prognosis and type of care of this patient population.

All LR*Efm* isolates were clearly LZD resistant by disk diffusion testing. MIC testing yielded a wide range of concentrations but those were not associated with the number of mutated 23S rRNA genes, in contrast to previous reports [4, 5]. TZD MICs have been previously shown to be lower than LZD MICs in *E. feacium*, which was also the case in our study [30]. However, the clinical significance of this is unknown, since clinical breakpoints have not yet been defined.

Interestingly, a genetic basis for the phenotypic LZD resistance could not be identified in all isolates. The only resistance mechanism identified in 14 (31.1%) of the LR*Efm* in our study was the G2576U point mutation in the 23S rRNA, meaning that additional resistance mechanisms must exist among Austrian LR*Efm* isolates, as proposed elsewhere [31]. According to a recent systematic review [32], up to 80.5% of LR*Efm* carry the point mutation G2576U,
although other authors have reported lower frequencies [10]. The previously undescribed additional point mutation A2598G is of unknown significance, however it was found only in three LREfm and only in combination with the G2576U mutation. The plasmid-mediated LZD resistance genes optrA, cfr and postA could not be detected by PCR nor by WGS.

Regarding resistance mechanisms apart from ribosomal point mutations, in a recent study on Mycobacterium abscessus [33], authors identified efflux pumps LmrS and mmpL9 at higher transcriptional levels among LZD-resistant isolates, suggesting an association with the resistant phenotype. Similarly, one could hypothesize that efflux pumps may contribute to LZD resistance, since efmA was detected among nearly all LREfm isolates. In addition, dfrF, which encodes for a chromosomal dihydrofolate reductase [34], was significantly associated (Fisher’s exact = 0.002) with the LREfm phenotype. Moreover, from a total of 11 cases that were treated with trimethoprim/sulfamethoxazole, eight cases belonged to the outbreak cluster. Trimethoprim is known to act against the reductase and sulfonamides have been described as a risk factor for LZD resistance acquisition [11].

Concerning the VGs, most of them (7/10) have been associated with more pathogenic E. faecium clones [25]. In accordance to other studies, sgrA, acm, ecbA are usually present at high frequencies in E. faecium strains of clinical origin [35, 36]. We did find an association between the adherence gene ecbA and the LREfm phenotype. Similarly, ecbA has been found in E. faecium outbreaks [37].

In our setting, the established control measures managed to terminate the initial outbreak event, although they did not prevent later sporadic cases. Unfortunately, environmental sampling was not performed and therefore an environmental reservoir was not identified. Nevertheless, the persistence of enterococci, especially of ST117, in hospital settings is well known [38] and as it was not identified for an environmental reservoir was not identified.

Conclusions
We have described an outbreak caused by Linezolid resistant Enterococcus faecium (LREfm) in an Austrian hematooncology unit. Whole-genome sequencing was a useful method to investigate isolates from the outbreak hospital and to compare them with strains from other healthcare facilities in Austria. The outbreak-causing strain was identified as highly clonal hospital adapted ST117, CT24 LREfm. Multiple additional clusters were identified, partly explaining the perceived repeated occurrence of LREfm after termination of the outbreak by multi-modal infection control measures. In a high-risk setting, active surveillance of LREfm is important for prompt outbreak detection.

Additional file

Additional file 1: Table S1. Resistance genes and point mutations at the 23S rRNA found among the sequenced strains. Table S2. Virulence genes carried by the 54 sequenced strains and resistance phenotype. (XLSX 17 kb)

Abbreviations
CC: clonal complex; cgMLST: Core genome multi locus sequence typing; CT: cluster type; DfM1: Department of Internal Medicine 1; EUCAST: European Committee on Antimicrobial Susceptibility Testing; LREfm: Linezolid resistant Enterococcus faecium; LSEfm: Linezolid susceptible Enterococcus faecium; LZD: Linezolid; ST: Sequence type; VAN: Vancomycin; VRE: Vancomycin resistant enterococci; WGS: Whole genome sequencing

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Authors’ contributions
HK and AC contributed equally and share first authorship. Study design: HK, AC. Epidemiological study: HK, AC, SMS. Microbiological analysis: HK, RH, PA. WGS: AC, WR. Data analysis: HK, AC, WR. Manuscript writing: HK, AC. Manuscript revision: FA, PA, WR. All authors read and approved the final manuscript.

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Availability of data and materials
The raw WGS reads were deposited into Sequence Read Archive (SRA) database under NCBI accession PRJNA541232. The datasets used and/or analyzed during the current study are either included in the published article and its supplementary files or available from the corresponding author on reasonable request.

Ethics approval and consent to participate
The study was approved by the local ethics committee (vote ECS 1053/2018).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.
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