for food safety rules and human health risk measures taken by national health and veterinary agencies. Regarding adequate processing of clinical samples and their preservation for morphologic and genetic evaluation, we strongly recommend fixation of positive fecal samples with eggs or segments (proglottids) immediately with 96%–99% molecular grade (i.e., not denatured) ethanol for future molecular diagnosis (1,4,8).

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Drug Resistance in Salmonella enterica ser. Typhimurium Bloodstream Infection, Malawi

To the Editor: Salmonella enterica serotype Typhimurium is one of the most common causes of bloodstream infection in sub-Saharan Africa (1). Among adults, the principal risk factor for invasive nontyphoidal Salmonella (iNTS) disease is advanced HIV infection; up to 44% of HIV-infected patients experience bacteremic recurrence through recrudescence of the original infection (2,3). Epidemics of iNTS disease in sub-Saharan Africa have been associated with a novel genotype of S. enterica ser. Typhimurium of multilocus sequence type (ST) 313 that is rarely seen outside the region and is associated with multidrug resistance (MDR) to chloramphenicol, cotrimoxazole, and ampicillin (4,5). As a consequence, ceftriaxone has become a key agent in the empirical management of nonfocal sepsis in Malawi (6).

In March 2009, a 40-year-old HIV-infected and antiretroviral therapy– naïve woman sought care in Blantyre, Malawi, with an MDR S. enterica ser. Typhimurium bloodstream infection. She was treated with ceftriaxone (2 g intravenously once daily) and discharged with oral ciprofloxacin (500 mg twice daily) for 10 days. She was readmitted 1 month later with recurrent fever. At this time, she had an MDR S. enterica ser. Typhimurium bloodstream infection with additional resistance to ceftriaxone and ciprofloxacin. In the absence of a locally available effective antimicrobial drug, she was treated with ceftriaxone, gentamicin, and high-dose ciprofloxacin but died shortly thereafter.

To help clarify how this extended MDR S. enterica ser. Typhimurium emerged, we determined the molecular mechanisms underpinning this disturbing pattern of antimicrobial resistance
We conducted phenotypic drug susceptibility testing by disk diffusion on *S. enterica* ser. Typhimurium strains A54285 (initial presentation) and A54560 (recurrence); both isolates were resistant to ampicillin, chloramphenicol, and cotrimoxazole, but A54560 exhibited additional resistance to ceftriaxone, ciprofloxacin, and tetracycline.

Paired-end sequencing of isolates A54285 (European Nucleotide Archive [ENA] accession number ERS035867) and A54560 (ENA accession no. ERS035866) that were cultured 1 month apart showed no differences between the conserved regions of these genomes (Figure). The similarity of these *S. enterica* ser. Typhimurium genomes strongly suggests that this recrudescence occurred after incomplete clearance of the first infection; although re-infection from the same source is unlikely, it cannot be excluded. Comparison of the accessory genomes, however, showed an additional 300 kb DNA in A54560.

Plasmid extraction and gel electrophoresis of genomic DNA identified a plasmid migrating in the gel to a position approximately equivalent to 120 kb, the size of ST313 virulence plasmid pSLT-BT in both strains, but no 300-kb plasmid was visualized in the ceftriaxone- and ciprofloxacin-resistant strain (A54560, data not shown), possibly because of the difficulty large plasmids have entering standard 1% agarose gels. However, ceftriaxone resistance was mobilized to *Escherichia coli* by conjugation at a frequency $6.5 \times 10^{-2}$ transconjugants per donor at 26°C. This frequency dropped dramatically to $\approx 1 \times 10^{-7}$ transconjugants per donor when conjugation was performed at 37°C. The presence of an IncHI2 plasmid in the transconjugants was confirmed by PCR for the IncHI2 region (7), and drug susceptibility testing confirmed that transconjugant clones acquired resistance to ceftriaxone, ciprofloxacin, and tetracycline.

These data confirm the presence of an extended-spectrum b-lactamase (ESBL)–producing IncHI2 plasmid in strain A54560 that is capable of conjugative transfer and suggest that the plasmid might have been acquired by residual index strain within the patient by transfer from an unknown donor.
bacterium. Partial decolonization of the patient’s gastrointestinal tract by ceftriaxone and fluoroquinolone antimicrobial therapy might have rendered it receptive to colonization by ESBL-producing bacteria, which we hypothesize donated the plasmid to the residual index strain.

The transconjugant plasmid DNA was sequenced by using the PacBio RSII platform (Pacific Biosciences, Menlo Park, CA, USA; http://www.pacificbiosciences.com), which assembled as a single contiguous sequence of 309,406 bp, designated pSTm-BTCR (online Technical Appendix Figure, ENA accession no. LK056646). We identified 331 predicted coding sequences, including 109 genes required for replication and transfer and 61 genes predicted to encode hypothetical genes. Fifteen genes associated with metabolism, membrane, virulence, antimicrobial resistance, and a toxin/antitoxin addiction system. We found an additional 160 predicted coding sequences, including 9 genes required for replication and transfer and 61 genes predicted to be associated with metabolism, membranes, virulence, antimicrobial resistance, and a toxin/antitoxin addiction system. We found an additional 160 predicted sequences, including 9 genes required for replication and transfer and 61 genes proposed to encode hypothetical genes.

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Drug Resistance in *Salmonella enterica* ser. Typhimurium Bloodstream Infection, Malawi

Technical Appendix

Methods

Bacteria

*Salmonella* Typhimurium A54285 (index multidrug-resistant) and A54560 (recurrence extended multidrug resistant) were isolated from the same patient at initial presentation and at recurrence 1 month later, respectively. Phenotypic drug susceptibility testing was undertaken by the disk diffusion method (Oxoid, Basingstoke, UK). The sequence of *Salmonella* Typhimurium D23580, an invasive isolate of multilocus sequence type ST313, isolated at Queen Elizabeth Central Hospital in 2004 (1), was used as a reference for sequence comparison. A rifampicin-resistant mutant of *Escherichia coli* C600 containing no plasmids was used as a recipient for conjugation.

Sequencing and Comparative Genome Analysis

Genomic DNA from bacteria was prepared by Wizard genomic DNA purification kit (Promega, Madison, WI, USA). Short-read Illumina sequencing libraries were prepared, and paired-end sequencing was performed on 2 μg gDNA on a HiSeq2000 platform (Illumina, San Diego, CA, USA) generating 150-bp reads (2). The whole-genome sequence of each isolate was assembled by using Velvet (http://www.ebi.ac.uk/~zerbino/velvet, version 1.2.03) and aligned against D23580 and its associated plasmids by using Abacas (version 1.3.1) (3). Single-nucleotide polymorphisms (SNPs) were determined with reference to strain D23580 by using SMALT, version 0.5.8 (Wellcome Trust Sanger Institute, Cambridge, UK). Contigs that failed to align with D23580 were considered as potentially unique mobile genetic elements and were further characterized by using BLASTn analysis (http://www.ncbi.nlm.nih.gov/BLAST) to predict whether they were plasmids; if so, in silico PCR was used to determine the plasmid incompatibility group (4).
Plasmid Isolation and DNA Extraction

Plasmid DNA was initially extracted by using an alkaline lysis method (Kado and Liu) and plasmids were profiled by visualising on a 1% agarose gel (5).

Individual plasmids were extracted following conjugal transfer from Salmonella Typhimurium A54560 to rifampicin-resistant E. coli K12 recipient strain c600, as described previously (6). Donor and recipient strains were mated by mixing 100 μL of overnight cultures on Luria-Bertani agar plates and incubated overnight at 26°C. Cells were cultured on MacConkey agar containing rifampicin (100 μg/mL) to select for E. coli and ceftriaxone (4 μg/mL) to select for transconjugants. The number of donor cells was determined by culture on Luria-Bertani agar containing rifampicin alone, and the transfer frequency was determined and the antimicrobial drug resistance phenotype of transconjugants was determined (Oxoid, USA), to ensure all resistances from the donor strain transferred.

Plasmid DNA was extracted from transconjugants by using an alkaline lysis method optimized for large plasmids, as previously described (7). PCR-based replicon typing on the transconjugant DNA was used to confirm the incompatibility group suggested by in silico plasmid typing (4).

Plasmid Sequencing

The transconjugant plasmid DNA was sequenced by using the PacBio RSII platform (Pacific Biosciences). Gene prediction and annotation was performed using Prokka (http://vicbioinformatics.com).

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Technical Appendix Figure. Plasmid map of pSTm-BTCR. Circles represent the 309-kbp circular plasmid with *rep1A* close to position 0. The outermost circle shows genes predicted on the forward strand, and the circle inside that shows genes predicted on the reverse strand. Red indicates genes involved in DNA replication; yellow indicates metabolism; purple indicates antimicrobial resistance or virulence; and green marks indicate hypothetical genes. Key resistance genes are annotated. The antimicrobial resistance genes are all located between 128 kbp and 182 kbp. The regions marked in blue in the next track highlights the sites of IS elements. The innermost circle shows the guanine–cytosine plot.