Horsing Around: Escherichia coli ST1250 of Equine Origin Harboring Epidemic IncHI1/ST9 Plasmid with bla_{CTX-M-1} and an Operon for Short-Chain Fructooligosaccharide Metabolism

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ABSTRACT  The relatedness of the equine-associated Escherichia coli strain ST1250 and its single- and double-locus variants (ST1250-SLV/DLV), obtained from horses in Europe, was studied by comparative genome analysis. A total of 54 isolates of E. coli ST1250 and ST1250-SLV/DLV from healthy and hospitalized horses across Europe (Czech Republic [n=23], The Netherlands [n=18], Germany [n=9], Denmark [n=3], and France [n=1]) from 2008 to 2017 were subjected to whole-genome sequencing. An additional 25 draft genome assemblies of E. coli ST1250 and ST1250-SLV/DLV were obtained from the public databases. The isolates were compared for genomic features, virulence genes, clade structure, and plasmid content. The complete nucleotide sequences of eight IncHI1/ST9 plasmids and one IncHI1/ST2 plasmid were obtained using long-read sequencing by PacBio or MinION. In the collection of 79 isolates, only 10 were phylogenetically close (≤8 single nucleotide polymorphisms [SNP]). The majority of isolates belonged to phylogroup B1 (73/79 [92.4%]) and carried bla_{CTX-M-1} (58/79 [73.4%]). The plasmid content of the isolates was dominated by IncHI1 ST9 (56/62 [90.3%]) and ST2 (6/62 [9.7%]), while 84.5% (49/58) of the bla_{CTX-M-1} genes were associated with the presence of the IncHI1 replicon of ST9 and 6.9% (4/58) with the IncHI1 replicon of ST2 within the corresponding isolates. The operon for the utilization of short-chain fructooligosaccharides (the fos operon) was present in 55 of 79 (69.6%) isolates, and all of these carried IncHI1/ST9 plasmids. The eight complete IncHI1/ST9 plasmid sequences showed the presence of bla_{CTX-M-1} and the fos operon within the same molecule. Sequences of IncHI1/ST9 plasmids were highly conserved (>98% similarity) regardless of country of origin and differed only in the structure and integration site of the multidrug resistance (MDR) region. E. coli ST1250 and ST1250-SLV/DLV are phylogenetically diverse strains associated with horses. A strong linkage of E. coli ST1250 with the epidemic multidrug resistance plasmid lineage IncHI1/ST9 carrying bla_{CTX-M-1} and the fos operon was identified.

KEYWORDS  Escherichia coli, IncHI1, ST1250, horses, multidrug resistance, plasmids
Extended-spectrum beta-lactamase (ESBL)-producing Enterobacteriaceae, such as ESBL-producing Escherichia coli (ESBL-PEC), are widespread in humans, food-producing and companion animals, wildlife, and the environment (1). ESBLs can be encoded by a variety of genes, among which the \(\text{bla}_{\text{CTX-M}}\) gene family dominates. The spread of \(\text{bla}_{\text{CTX-M}}\) in ESBL-PEC is facilitated by mobile genetic elements, including multidrug resistance (MDR) plasmids of various incompatibility (Inc) groups (2, 3).

Antimicrobial resistance is of concern for a wide range of equine pathogens. ESBL-producing Enterobacteriaceae have been increasingly isolated from wounds, as well as from individuals with respiratory and urinary tract infections (4). Several studies have reported the occurrence of CTX-M-1-producing \(E. \text{coli}\) in hospitalized but also nonhospitalized horses in many European countries, such as the Czech Republic (5, 6), Denmark (7, 8), The Netherlands (9, 10), the United Kingdom (11), Germany (12, 13), Sweden (14), Switzerland, and France (14), including Guadeloupe (15).

An epidemic IncHI1/ST9 plasmid lineage conferring multidrug resistance and harboring \(\text{bla}_{\text{CTX-M-1}}\) has been identified in \(E. \text{coli}\) isolates from hospitalized horses in the Czech Republic (5) and later in isolates from healthy and diseased horses in France (16), Sweden (14), and The Netherlands (10). Moreover, the \(\text{fos}\) operon, consisting of seven genes (\(\text{fosK, fosY, fosGH2, fosX, fosGH1, fosT, fosR}\)) involved in the metabolism of short-chain fructooligosaccharides (scFOS), has been detected on pEQ1, a conjugative IncHI1/ST9 plasmid obtained from \(E. \text{coli}\) T23, originating from the feces of a hospitalized horse in the Czech Republic (6). The importance of the presence of the \(\text{fos}\) operon arises from the fact that until Schouler et al. (17) detected the \(\text{fos}\) operon in extraintestinal pathogenic \(E. \text{coli}\), the \(\text{fos}\) operon occurred only in probiotic, not pathogenic, bacteria and only as part of a chromosomally encoded genomic island. However, none of the studies mentioned above, except for that of Dolejska et al. (6), characterize the complete sequence of an epidemic IncHI1/ST9 plasmid. The presence of the \(\text{fos}\) operon might play a role in the coselection of multiresistance plasmids, since it provides bacteria with a metabolic advantage and enables them to grow in environments where pathogenic bacteria would normally not survive. The association of the \(\text{bla}_{\text{CTX-M-1}}\) gene and the \(\text{fos}\) operon on plasmids is particularly worrying because of the specific role of fructooligosaccharides in the horse diet and the ability of plasmids to be transferred horizontally. scFOS are commonly used prebiotics, and they are often part of a supplementary horse diet which should support gut health and beneficial microflora in horses (18). However, our results indicate that using scFOS in a horse diet may contribute to the spread of antibiotic resistance genes, so it is doubtful whether scFOS are indeed beneficial. Moreover, the use of scFOS as prebiotics is not limited to horses; they are part of many human probiotics and nutritional supplements for newborns. Therefore, more attention should be paid to the relation between the ability to utilize scFOS and antibiotic resistance.

In the current study and publicly available sequence data, \(E. \text{coli}\) ST1250 is found to be a rare sequence type (ST) in hosts other than horses. Moreover, the close association of \(E. \text{coli}\) ST1250 with IncHI1/ST9 has been observed (10, 14, 16). Clinical and nonclinical isolates of \(E. \text{coli}\) ST1250 and its single- and double-locus variants (ST1250-SLV/DLV) collected from horses in the Czech Republic, The Netherlands, Denmark, France, and Germany were subjected to short-read DNA sequencing. \(E. \text{coli}\) ST1250 was selected due to its frequent occurrence in horse samples. Other STs included are single- or double-locus variants of ST1250 and were selected both for their close relationship (with regard to the multilocus sequence type [MLST]) to ST1250 and for their frequent occurrence in horse samples. For the selected isolates, long-read sequencing was applied to obtain complete nucleotide sequences of chromosomes and IncHI1 plasmids. Additional sequences of \(E. \text{coli}\) ST1250 and ST1250-SLV/DLV were recovered from EnteroBase before genomic features, clade structure, and plasmid content were examined by detailed phylogenetics. Our hypothesis is that \(E. \text{coli}\) ST1250 is adapted to horses and has low pathogenic potential itself, while this study shows frequent
association of *E. coli* ST1250 with HI1/ST9 multiresistance plasmids encoding *bla*$_{CTX-M-1}$ and the *fos* operon.

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**RESULTS AND DISCUSSION**

**Genomic characterization of the isolates.** A total of 79 draft genomes of *E. coli* ST1250 or ST1250-SLV/DLV, consisting of 54 sequences obtained in this study and 25 retrieved from EnteroBase or GenBank, were analyzed. The majority of isolates belonged to phylogroup B1 (73/79 [92.4%]), followed by phylogroups A (5/79 [6.3%]) and E (1/79 [1.2%]). Since the collection studied was aimed at *E. coli* ST1250 and ST1250-SLV/DLV, the majority of the isolates in the collection indeed belonged to ST1250 (60/79 [75.9%]), followed by ST1250-SLV/DLV: DLV-ST4164 (4/79 [5.0%]), SLV-ST826 (3/79 [3.3%]), SLV-ST4527 (2/79 [2.5%]), SLV-ST1686 (2/79 [2.5%]), SLV-ST11123 (2/79 [2.5%]), and SLV-ST2336, SLV-ST4552, SLV-ST7434, DLV-ST11033, SLV-ST11121, and SLV-ST11122 (1/79 [1.2%] each). The common detection of *E. coli* ST1250, along with ST1250-SLV/DLV, in horses as opposed to other sources suggests that this ST is horse associated. This assumption is supported by our unpublished study of hospitalized and healthy horses across the Czech Republic, where 31.7% (20/63) of *E. coli* isolates belonged to ST1250 (6/63 [9.5%]) or ST1250-SLV/DLV (14/63 [22.2%]). Similarly, in a nationwide study conducted on nonhospitalized horses from The Netherlands (10), the most prevalent group of isolates whose whole genomes had been sequenced belonged to ST1250 (8/48 [16.7%]) or ST1250-SLV/DLV (9/48 [18.8%]). Isolates from both the Czech Republic (unpublished data) and The Netherlands are included in this study. Despite the limited number of publicly available draft genome assemblies of *E. coli* ST1250, the association of *E. coli* ST1250 with horses (7/10 [70%]) was observed in EnteroBase as well, where only 1 of 10 ST1250 isolates was of human origin, while 2 were of undefined origin.

Antibiotic susceptibility testing of the isolates that were physically available (*n* = 54) showed that all exhibited a multidrug (≥3 antibiotics) resistance profile corresponding to a resistance genotype, except for one (d16-1) that was resistant to ciprofloxacin but susceptible to nalidixic acid, yet no *qnr* gene was detected. Among the 79 sequences analyzed for antibiotic resistance gene (ARG) content, the most prevalent ESBL genes included *bla*$_{CTX-M-1}$ (58/79 [73.4%]), *bla*$_{CTX-M-2}$ (6/79 [7.6%]), and *bla*$_{SHV-12}$ (7/79 [8.9%]). While 4 isolates carried both *bla*$_{CTX-M-1}$ and *bla*$_{SHV-12}$, 12 isolates did not encode any ESBL. The whole-genome sequence (WGS) data from 79 isolates showed the presence of genes conferring resistance to tetracycline, trimethoprim, sulfonamides, gentamicin, streptomycin, and chloramphenicol (see Fig. S1 in the supplemental material). Neither the resistance profile nor the plasmid content was influenced by the source of isolation (feces or clinical material). A complete profile of ARGs, along with the results of antibiotic susceptibility testing, can be found in Fig. S1.

Analysis of virulence genes showed the presence of the siderophore synthase-encoding operon *ent* and the enterobactin export-encoding gene *fes* in all 79 isolates and the enterobactin uptake-encoding operon *fep* in 75 (94.9%) isolates. These genes are common for the majority of Gram-negative bacteria (19) and may be considered beneficial for the animal hosting the bacteria carrying these genes. This has been observed in *Caenorhabditis elegans*, where the bacterial enterobactin promoted the growth of *C. elegans* via mitochondrial iron uptake (20). More than half of the isolates (56/79 [70.9%]) harbored the *fli* operon, which is involved in the biosynthesis and functioning of the flagellar organelle (21). Twelve of the 79 isolates (15.2%) carried a *fim* operon, encoding type I fimbriae. The results of *in silico* screening for virulence genes suggest that the isolates did not carry any virulence factors enabling their classification into *E. coli* pathotypes. These findings support the observation that *E. coli* ST1250 and related STs are part of the commensal equine microbiota (10).
Regarding the plasmid content, the most common plasmid replicons among the collection of 79 isolates were three IncHI1-related replicons (62/79 [78.5%]) of ST9 (56/62 [90.3%]) and ST2 (6/62 [9.7%]). The high prevalence of conjugative IncHI1/ST9 and its association with \( \text{bla}_{\text{CTX-M-1}} \) have been reported recently (16). IncHI1/ST9 appears to be very well adapted to \( E. coli \) in horses, especially \( E. coli \) ST1250.

**Comparison of \( E. coli \) ST1250 and \( E. coli \) K-12 chromosomes.** The results revealed that unlike \( E. coli \) K-12, the majority of \( E. coli \) ST1250 and ST1250-SLV/DLV strains did not carry genes encoding products facilitating adhesion and colonization (\( \text{yadN, ygiL, paa, yde} \)) (22, 23), a type II secretion pathway (\( \text{gspCO} \)) (24, 25), the stress response (\( \text{ygfK, yqeK, ydfO} \)) (26–28), short-chain fatty acid and xanthosine metabolism (\( \text{ato, xapA} \)) (29, 30), or a toxin-antitoxin system (\( \text{rnlAB} \)) (31). This finding suggests that \( E. coli \) ST1250 does not contain many virulence genes and represents a lineage with low pathogenicity that is adapted to the horse gut. Generally, in terms of genetic features that are conserved and specific for \( E. coli \) ST1250, no per se pathogenic genes were detected in the genome.

**High diversity of ST1250 and ST1250-SLV/DLV.** Phylogenetic analysis based on the core genome from whole-genome sequences of 79 isolates showed high diversity of \( E. coli \) ST1250 and ST1250-SLV/DLV (Fig. 1; also Fig. S1). The core genome alignment was created from 2,556 core genes and had a length of 3,382,416 bp. The isolates differed in 0 to 10,490 single nucleotide polymorphisms (SNP), with 10 phylogenetically close isolates differing in only 0 to 8 SNP.

The isolates in the minimum spanning tree were clustered regardless of their geographical origin and the carriage of IncHI1 plasmids (Fig. S2). Despite the high variability, three noteworthy clusters with one subcluster, marked A, B1, BII, and C, emerged (Fig. 1). Cluster A consisted of six ST1250-SLV/DLV isolates, including ST4164 (four isolates) and ST4527 (two isolates), from The Netherlands, the Czech Republic, and Germany. Their SNP differences ranged from 29 to 321, while the lowest SNP difference (29 SNP) was between a Dutch and a German isolate (15S04665-1 and IHIT_19665). Cluster B was made of subclusters B1 and BII, consisting of three and four Czech ST1250 isolates with SNP differences of 1 to 16 and 0 to 8, respectively. Cluster C consisted of nine ST1250 isolates and one novel SLV. The isolates in this cluster differed in 0 to 530 SNP and originated from Switzerland, The Netherlands, the Czech Republic, Denmark, and Germany. However, presumed clonal spread of some isolates occurred only sporadically, usually within the same country. For example, no SNP differences in cluster C were observed between two couples of Swiss isolates (ESC_JA8778AA with ESC_ZA2274AA and ESC_JA8764AA with ESC_ZA2276AA).

Our findings suggest that even though there are clusters (A and C) of closely related international isolates, the isolates in these clusters belonged to several different serotypes (Fig. 1), and no clonal spread (<10 SNP) of \( E. coli \) ST1250 was demonstrated. Closely related isolates were observed only within subclusters B1 and BII, which consist largely of isolates from the same country, sampling site, and year. Subcluster B1 consisted of three isolates obtained within the same horseback riding center in 2015. Subcluster BII consisted of four isolates, of which three were obtained from a horse clinic in 2008 and one (isolate 97989) from a horseback riding center in 2015. Isolate 97989 was closely related (8 SNP) to the three isolates obtained from the horseback riding center. However, it is not clear whether this relatedness is caused by the persistence of the clone within the sampling sites over several years or by repeated introduction of the clone to the horse outside the hospital.

Phylogenetic analysis of our international collection of \( E. coli \) ST1250 isolates, including ST1250-SLV/DLV, indicates that \( E. coli \) ST1250 is genetically highly variable yet apparently associated with horses, since to our knowledge, this ST was not commonly detected in nonequine studies. However, there is no evidence of transmission of ESBL-Pec ST1250 to humans, except for one record of \( E. coli \) ST1250 of human origin in EnteroBase, with no further information. Moreover, \( E. coli \) ST1250-SLV/DLV are closely related to \( E. coli \) ST1250 clonal lineages, highlighting the importance of whole-
FIG 1 Phylogenetic tree depicting the diversity of E. coli ST1250 and ST1250-SLV/DLV isolates with highlighted clusters A, BI, BII, and C. Isolates marked *M or *PB were sequenced on MinION or PacBio, respectively. Metadata for country of origin, year of isolation, source, ST, phylogroup, and serotype are included. Colored boxes indicate the presence of the IncHI1 plasmid and its ST, the \textit{fos} operon, and ESBL genes. The designations of physically obtained isolates are set in boldface, and data sets from EnteroBase start with "ESC."
Comparative genomics of IncHI1 plasmids. IncHI1 plasmids were identified in the genomes of 62 E. coli strains in this study. Comparison of draft genomes and complete IncHI1 plasmids revealed two lineages of IncHI1: ST2 (6/62 [9.7%]) and ST9 (56/62 [90.3%]). These lineages are distinguished by one SNP in repHIA (allele HCM1_064), plasmid nucleotide identity (between p10068 [IncHI1/ST2] and p99063 [IncHI1/ST9]) of >99.9%, and >96% query coverage. They differ in the structure of the multidrug resistance (MDR) region and by the presence of the fos operon in IncHI1/ST9 plasmids. Once these regions were removed, comparison of the plasmid backbones of p10068 and p99063 showed nucleotide identity and query coverage of 99.98% and 99%, respectively.

The majority (45/56 [80.4%]) of isolates with IncHI1/ST9 plasmids harbored a class 1 integron with a dfrA17-aadA5 gene cassette. Isolates carrying IncHI1/ST2 plasmids (5/6 [83.3%]) were associated with a class 1 integron with a dfrA12-aadA2 cassette array. The other genetic features of the isolates harboring IncHI1/ST2 and IncHI1/ST9 plasmids are described in the figures presented below and in the supplemental materials. Eight complete nucleotide sequences of IncHI1/ST9 were obtained using PacBio (n = 4) and MinION (n = 4) long-read sequencing. Overall, they exhibited high nucleotide identities (>98%) to pEQ1 and to each other. Figure 2 shows a comparison of complete and draft genome assemblies of IncHI1/ST9 plasmids. Two different variants of the integration of an MDR region were identified within the eight complete plasmid sequences. Five plasmids harbored an MDR region of 41,754 bp (p15S04714-1) to 49,700 bp (p15S0829-4) between genes encoding a hypothetical protein of the backbone (Fig. 3, p15S04714-1). The very same position of the MDR region is found in pEQ1, of equine origin, and pHCM1 (GenBank accession no. AL513383), of human origin, designated a type 2 IncHI1 plasmid based on integration sites of the MDR region inside the plasmid backbone (32). The other three plasmids contained an MDR region divided into two parts (14,708 bp and 34,143 bp) by a part of the inverted plasmid backbone of 14,228 bp (Fig. 3, p99063). This inversion was presumably mediated by two copies of IS26 flanking the backbone part (14,228 bp) and the larger, 34,143-bp part of the MDR region. This finding points to the existence of two variants of highly conserved epidemic IncHI1/ST9 plasmids differing in genetic structure, length, and the integration site of the MDR region. Besides blacTX-M-1, each of the eight complete IncHI1/ST9 plasmids carried genes encoding resistance to tetracycline [tet(B)], trimethoprim (dfrA17), gentamicin [aac(3)-Ild], sulfonamides (sul1, sul2), and streptomycin [aph(6)-Ia, aph(3’)-Ib, aadA5] in the MDR region. However, they differed in the presence of the mph(A), aph(3’)-Ia, catA1, and blalTEM-1 genes. A Mauve comparison of the MDR region variants and their rearrangements is shown in Fig. S3. The mer operon, qacEΔ1 gene, and chrA gene, encoding determinants for resistance to mercury, quaternary ammonium compounds, and chromium, respectively, were present in each complete IncHI1/ST9 plasmid, too (Fig. 2).

The MDR region of the complete IncHI1/ST2 plasmid obtained using MinION (p10068) was integrated between genes encoding a hypothetical protein of the backbone, identical to the integration site in IncHI1/ST9 type 2 plasmids. Unlike IncHI1/ST9 plasmids, IncHI1/ST2 plasmids did not harbor the fos operon. Since only one complete IncHI1/ST2 plasmid was obtained, the identity of p10068 could not be properly assessed in comparison with the draft genome assemblies from other IncHI1/ST2 plasmids (n = 5) in this study. However, a BLAST Ring Image Generator (BRIG) comparison of p10068 with the draft genome assemblies of the other five IncHI1/ST2 plasmids (Fig. S4) showed low variability in the backbone and the MDR region of IncHI1/ST2 plasmids from the Czech Republic, Canada, and Switzerland, pointing toward the geographically widespread potential of the IncHI1/ST2 plasmid lineage. Plasmid p10068 harbored the blacTX-M-1, tet(B), sul1, sul2, aac(3)-Ild, aadA2, dfrA12, aph(6)-Ia, and aph(3’)-Ib genes within the 40,759-bp MDR region.
Even though IncHI1/ST9 and IncHI1/ST2 plasmids have highly conserved backbones and differ mostly within the MDR region, IncHI1/ST2 plasmids did not harbor the *fos* operon. A higher prevalence of IncHI1/ST2 over IncHI1/ST9 has been observed only in Swedish isolates so far (14); however, no data on the presence of the *fos* operon are available from that study. In summary, comparison of IncHI1/ST9 plasmids of various geographical origins showed a highly conserved structure and high nucleotide identity, highlighting the epidemic potential of IncHI1/ST9 plasmids. Similarly, plasmid lineage IncI1-h, harboring Ambler class A and C genes from *E. coli* and *Salmonella enterica* of animal and human origins, has been found to be epidemic (33), and IncX3 plasmids encoding NDM-type enzymes within different enterobacterial species (34) or specific NDM-5-encoding IncX3 plasmids from *E. coli* strains of diverse sequence types have been obtained from five different pig farms from different geographical regions of China (35).

The *fos* operon is highly diffused among ST1250 strains and is carried by IncHI1/ST9 plasmids. The collection of 79 *E. coli* ST1250 and ST1250-SLV/DLV isolates was screened in silico for the presence of the 8,896-bp *fos* operon, which is involved in the metabolism of scFOS. Fifty-eight (58/79 [73.4%]) isolates harbored the *fos* operon,
while 1 isolate (ESC_ZA2274AA_AS) lacked the fosK gene, suggesting the presence of a nonfunctional operon. The majority (56/58 [96.6%]) of isolates harboring the fos operon also harbored an IncHI1/ST9 plasmid. A BRIG comparison including all IncHI1/ST9 draft genome assemblies from short-read sequencing data (n = 56) showed that the fos operon is highly conserved and is carried by this plasmid lineage (Fig. 2). The association of the fos operon with IncHI1/ST9 plasmids was first observed in equine isolates from the Czech Republic (6) and subsequently in The Netherlands (10) and France (16).

The broader genetic surrounding of the fos operon was examined within eight complete IncHI1/ST9 plasmids obtained using long-read sequencing. In all plasmids, the fos operon was conserved with 100% identity and surrounded by IS903B. As observed previously in pEQ1, the operon was part of a 24-kb module inserted between the repE (FIA replicon) and hns genes of the plasmid backbone (6). Only one of eight plasmids (p15S04779-4) differed within the 24-kb module; it harbored IS1203 upstream of the fos operon.

Although the association of the fos operon and the IncHI1/ST9 plasmid was evident, the fos operons of two E. coli ST1250 isolates (99783-3RV and T20) had different genetic contexts. Analysis of the genetic surroundings of the fos operon of the 99783-3RV isolate showed the presence of an IncFIB(K) replicon on the same contig, suggesting the carriage of the fos operon by plasmids other than IncHI1/ST9 plasmids. The limited information provided by the short-read sequencing did not enable us to identify the genetic context of the fos operon within the T20 strain. However, plasmid replicons IncFIB(K) and IncY were identified in T20, suggesting the possibility that the fos operon is either mediated by plasmid types other than IncHI1/ST9, as in the 99783-3RV isolate, or chromosomally encoded, as observed previously for E. coli BEN2908 (GenBank accession no. AY857617) (36).

It has been suggested that a plasmid-mediated sugar metabolic element, such as the fos operon, could play a key role in strain fitness, contributing to the successful dissemination and maintenance of the fos operon-containing MDR IncHI1/ST9 plasmids in the equine intestinal microbiota (6). We suppose that the absence of the fos operon in IncHI1/ST2 plasmids might contribute to their generally low prevalence, in contrast to IncHI1/ST9 plasmids, among equine E. coli isolates, as observed in our study, since the fos operon seems to provide isolates with a selective advantage in the presence of scFOS. The equine diet is usually rich in scFOS, which may, along with the usage of antibiotics, provide positive selective pressure for bacteria carrying this fos operon-associated plasmid lineage. However, the question of whether factors such as different amounts of scFOS in the horse feed or the geographical origin of the isolate are influencing the selection of the IncHI1/ST9 plasmid lineage needs to be clarified by further studies.

Interestingly, the presence of the fos operon in E. coli BEN2908 was linked to an increased ability to colonize the avian intestine (17). We hypothesize that the widespread
presence of the *fos* operon in *E. coli* ST1250 and ST1250-SLV/DLV isolates, as observed in our study, may compensate for the absence of known chromosomally encoded colonization factors in these strains. Further elucidation of the role of the *fos* operon in the colonization of the horse intestine by *E. coli* ST1250 would require in vivo experiments.

The genomic study of *E. coli* ST1250 and ST1250-SLV/DLV isolates of equine origin has shown great chromosomal diversity, which could not be linked to specific geographic regions. In comparison, the highly prevalent ESBL-encoding IncHI1/ST9 plasmids in these strains show little diversity beyond the presence of specific MDR regions and the *fos* operon. To summarize, *E. coli* ST1250 and *E. coli* ST1250-SLV/DLV isolates seem to be well adapted to horses as hosts, and *E. coli* ST1250 in particular often shows an association with IncHI1/ST9 multiresistance plasmids encoding CTX-M-type enzymes and harboring the *fos* operon, which may contribute to the coselection of these plasmids.

**MATERIALS AND METHODS**

**E. coli isolates.** A total of 54 ST1250 and ST1250-SLV/DLV isolates from healthy and hospitalized horses in several countries across Europe from 2008 to 2017 were collected (Fig. S1 in the supplemental material, isolates in boldface). The isolates originated from the Czech Republic (*n* = 23) (5; unpublished data), The Netherlands (*n* = 18) (9), Germany (*n* = 9), Denmark (*n* = 3) (7), and France (*n* = 1) (14). Forty-two and 12 isolates were obtained by selective cultivation on MacConkey agar (MCA) with cefotaxime (1 or 2 mg/liter) and on MCA without antibiotics, respectively. The complete details regarding processing of the samples can be found in the corresponding studies mentioned above. The majority of the isolates originated from feces (42/54), various clinical materials (10/54), and environmental swabs (2/54) (Fig. S1). Susceptibility to 22 antibiotics (ampicillin [AMP], streptomycin [S], sulfonamides [S3], tetracycline [TE], trimethoprim-sulfamethoxazole [SXT], chloramphenicol [C], cefazolin [KZ], nalidixic acid [NA], ceftazidime [CAZ], gentamicin [CN], amoxicillin-clavulanic acid [AMC], ciprofloxacin [CIP], fosfomycin [FOT], erythromycin [ETP], imipenem [IMP], meropenem [MEM], aztreonam [ATM], nitrofurantoin [F], azithromycin [AZM], colistin [TC], ticarcillin [TGC], cefalothin [KF]) was tested by disk diffusion according to the CLSI (37), and the ESBL phenotype was determined by the MASTDISCS test (D68C1 AmpC & ESBL detection set; Mast Diagnostics, UK).

**Short-read sequencing.** All isolates were subjected to whole-genome sequencing (WGS) with at least 50× expected coverage. Genomic DNA was extracted using the NucleoSpin Tissue kit (Macherey-Nagel, Germany) and was used for DNA library preparation with a Nextera XT Library Preparation kit. Sequencing was performed on a MiSeq system (Illumina) using 2 × 250-bp paired-end sequencing.

**Publicly available data.** We searched Enterobase (http://enterobase.warwick.ac.uk/species/index/ecoli) for sequence data for *E. coli* ST1250 and ST1250-SLV/DLV, and assemblies along with corresponding metadata were downloaded. Available data sets for ST1250 isolates irrespective of the plasmid content and ST1250-SLV/DLV variants harboring IncHI1 were selected. Thirteen draft genome assemblies originating in Switzerland (*n* = 7), Germany (*n* = 2), Canada (*n* = 1), the United States (*n* = 1), and unknown origin (*n* = 2) were retrieved from Enterobase. Additional ST1250 and ST1250-SLV/DLV sequences from horses in The Netherlands (*n* = 11) and Guadeloupe (*n* = 1) were retrieved from BioProject (record numbers PRJEB34847 [10] and PRJNA383339 [15], respectively). In total, 25 draft genome assemblies of *E. coli* ST1250 and ST1250-SLV/DLV isolates were obtained from the public databases; among these, 22 originated from horses, 2 were of unknown origin, and 1 isolate was of human origin (Fig. S1).

**Long-read sequencing.** PacBio sequencing was used to obtain complete high-quality reference chromosomal and plasmid sequences. Subsequently, sequences of randomly selected IncHI1 plasmids were obtained using MinION sequencing. For PacBio sequencing, genomic DNA was extracted using a NucleoSpin Microbial DNA kit (Macherey-Nagel, Germany). Libraries were prepared according to the microbial multiplexing protocol. Shearing was performed using g-tubes (Covaris, USA), and no size selection was done during library preparation. Sequencing was performed with the Sequel I platform (Pacific Biosciences, Menlo Park, CA, USA). For Oxford Nanopore Technologies (ONT) sequencing, the DNA was extracted using a Gentra Puregene Yeast/Bact. kit and a DNeasy blood and tissue kit (Qiagen, Hilden, Germany). Libraries were prepared using a 1D ligation barcoding kit (catalog no. SQK-LSK109 and EXP-NBD104; ONT, Oxford, UK) and were sequenced in R9.4.1 and Flongle flow cells with a MinION Mk1B (ONT).

**Sequence data analysis.** Raw data from short-read sequencing were adaptor and quality (*Q* ≥ 20) trimmed using Trimmomatic (38), *v* 0.36. *De novo* assembly was performed using SPAdes (39), *v* 3.13.1, with the “careful” option. For PacBio data, HGAP4 (40) in SMRT Link, *v* 6.6, was used to perform the assemblies with a minimum seed coverage of 30. Long reads from MinION were demultiplexed and converted from fast5 to fastq format using Guppy, *v* 3.2.1 (ONT). The adaptors and low-quality ends (*Q* ≤ 8) were trimmed using Porechop, *v* 0.2.2 (https://github.com/rrwick/Porechop) and BBduk (https://sourceforge.net/projects/bbmap/), respectively. *De novo* hybrid assembly using short reads from Illumina and long reads from MinION was performed with Unicycler (41), *v* 0.4.7. The complete chromosomal and plasmid sequences were annotated using RASTtk (42) and were manually curated according to GenBank in Geneious, *v* 9.0.5.
Genomic characterization. The draft genome assemblies from WGS and EnteroBase were analyzed for the presence of antibiotic resistance genes (ARGs), virulence genes, plasmid replicons, and serotypes. For this purpose, ABRicate (https://github.com/tseemann/abricate) was used with thresholds set to 90% identity and 95% query coverage against the databases ResFinder (43), VFDB (44), PlasmidFinder (45), and EcOH (46), respectively. Plasmid multilocus sequence typing (pMLST) was performed using pMLST by the Center for Genomic Epidemiology (45). MLST profiles were verified by mlst (https://github.com/tseemann/mlst), and the phylogroups were assessed using ClermonTyping (47).

Phylogenetic analysis. The phylogenetic relatedness of isolates was evaluated. The open reading frames were predicted from draft genome assemblies using Prokka (48), and GFF3 files were used for core genome alignment in Roary (49) with default settings (the percentage of isolates a gene must be in to be considered a core gene is 99%). The alignment was then used in RaXML (50) for calculating the phylogenetic tree under the GTR+GAMMA model supported by 1,000 bootstraps. The tree was visualized in iTOL, v4 (51), and GrapeTree (52). The nucleotide similarity between the isolates was determined by assessing the number of single nucleotide polymorphisms using snp-dists (https://github.com/tseemann/snp-dist) with the core genome alignment as an input.

Comparative genomics. Comparison of four high-quality annotated complete chromosomal E. coli ST1250 sequences obtained from PacBio sequencing with E. coli K-12 as a reference (GenBank accession no. U00096) was performed using progressiveMauve (53). E. coli K-12 was selected especially because of the lack of virulence genes and the high level of characterization its genetic content. Therefore, the comparison allowed us to see if there are virulence factors or any other genes that could offer an advantage for E. coli ST1250, since E. coli K-12 is a nonpathogenic strain. The complete annotated plasmid sequences were compared with respect to their plasmid replicon and plasmid ST using BLAST Ring Image Generator (54). The structure of the fos operon (nucleotides 61904 to 70800 of pEQ1 [GenBank accession no. KF362121]) was examined and compared to the sequence within pEQ1.

Data availability. The draft genome assemblies, the complete plasmid and chromosomal sequences, and the raw data from short-read sequencing have been deposited in GenBank and SRA, respectively, under BioProject number PRJNA638007. The complete nucleotide sequences of the plasmids have been deposited in GenBank under accession numbers MT586601 to MT586609.

SUPPLEMENTAL MATERIAL
Supplemental material is available online only.

SUPPLEMENTAL FILE 1, PDF file, 8.3 MB.

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We have no potential conflicts of interest to declare.

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