

## Detailed Materials and Methods

**Subject selection and sample collection.** The study subjects were recruited among the subjects of a larger study in which Swedish students from the universities in Umeå, Stockholm and Gothenburg travelling for international exchange studies as part of their university education were invited to participate from April 2010 through January 2014. The study protocol included responding to questionnaire data and self-submission of fecal specimens before and after travel. Each study subject submitted a fecal swab specimen (Copan Venturi Transystem®, Copan diagnostics Inc. USA) for bacterial culture and a fecal specimen in a half filled 101\*16,5mm plastic collection tube (Faeces tube, Sarstedt, Nümbrecht, Germany) for metagenomic analysis. For the present study, only healthcare students (medical, nursing and dentistry) travelling to the Indian peninsula or to Central Africa and having submitted a full set of fecal specimens qualified for inclusion. None of the subjects included were allowed to have taken antibiotics within six months prior to sampling. The inclusion was consecutive with a target of 15-20 study subjects travelling to each of the two destinations.

Fecal specimens for metagenomics were sent to the lab in the collection tubes, and stored at -20°C upon arrival. DNA extraction was performed using the QIAamp® DNA Stool Mini Kit (QIAGEN) according to the manufacturer's protocol. DNA concentrations were measured using a Qubit instrument, and samples were stored at -20°C before sequencing. Paired-end sequencing libraries (2x100 bp) were prepared using the TrueSeq DNA Kit for multiplexing. Sequencing was done at Science for Life Laboratories (Stockholm, Sweden) using Illumina HiSeq2000 technology. The study was approved by the regional ethical review board in Umeå, Sweden (2011-357-32M).

**Culturing of ESBL-producing bacteria.** Screening for ESBL producing bacteria was done using swab culture on chromogenic media (crohmID ESBL, bioMérieux SA, Marcy-l'Etoile France). All positive isolates were analyzed through culture-based methods according to EUCAST guidelines (<http://www.eucast.org>). The definition of ESBL producing Enterobacteriaceae by Giske *et al.* (1) was used. Antibiotic susceptibility testing (including cefotaxime, ceftazidime, piperacillin/tazobactam and meropenem) was done by disc diffusion (Oxoid Ltd./Thermo Fisher Scientific, Cambridge, United Kingdom) on Mueller-Hinton (MH) agar (Oxoid Ltd.). E-tests® (bioMérieux) were used to test for the presence of ESBL<sub>A</sub> phenotype (detecting the presence of CTX-M, SHV and TEM enzymes) with cefotaxime/cefotaxime + clavulanic acid and ceftazidime/ceftazidime + clavulanic acid as well as cefepime/cefepime+ clavulanic acid if warranted. If applicable, phenotypic detection of AMPc type beta-lactamases with cloxacillin inhibitable ESBL enzymes (cefotetan/cefotetan + cloxacillin) was performed. Since not all carbapenemase-producers show reduced susceptibility to carbapenems, particularly not OXA-48/OXA-181 producers that do not yield an ESBL (2, 3), all post-travel samples of this study (before screening on chromogenic media) were tested by disc diffusion for susceptibility to meropenem (10µg), piperacillin/tazobactam (30µg/6µg) and ceftazidim (10µg) on MH agar (Thermo Scientific) containing 7,5mg/L vancomycin, a modified zone diameter cut-off ( $\geq 16$  mm) for piperacillin-tazobactam was used (4). Resistant Enterobacteriaceae isolates were further tested with temocillin (30 µg, Rosco Diagnostica, Taastrup, Denmark) and our test scheme contained additional analysis by the CT-103 XL microarray (Check-Points Health B.V., Wageningen, The Netherlands) of isolates with reduced sensitivity to temocillin. Phenotypic species identification of ESBL positive Enterobacteriaceae isolates was done using an API® identification system (API-20E, bioMérieux). Species identification of non-Enterobacteriaceae isolates growing on the chromogenic media was performed with MALDI-TOF-MS using a Bruker Daltonics Microflex

LT mass spectrometer and the MALDI-Biotyper 3.0 software (Bruker Daltonik GmbH, Bremen, Germany).

DNA from ESBL-producing strains was extracted using QIAamp® DNA Mini Kit (QIAGEN) according to the manufacturer's protocol. DNA concentrations were measured using a Qubit instrument, and samples were stored at -20°C before sequencing. The samples were sequenced using the Illumina MiSeq instrument (FOI, Umeå), producing 299,202 – 1,575,161 (average 760,896) 250 bp paired-end reads per sample for each library.

**Sequence pre-processing and quality filtering.** FASTQ sequences from both genomes and metagenomes were trimmed with respect to adapters and low-quality nucleotide stretches using TrimGalore! version 0.2.8, [www.bioinformatics.babraham.ac.uk/projects/trim\\_galore/](http://www.bioinformatics.babraham.ac.uk/projects/trim_galore/) with parameter settings “-stringency 6 -q 28”. Host sequences were subsequently removed from the metagenomic data by aligning the trimmed sequences to the hg19 human genome reference assembly using Bowtie2 v. 2.0.2 (5) with parameters “-p 8 -un-conc-gz *OUTFILE*” and discarding those reads that had a valid alignment to it. Trimmed and filtered sequences were used as input for all further analysis, including resistance gene mapping, reference-genome mapping, and *de novo* assembly. Sequence data have been submitted to the European Nucleotide Archive under project accession number PRJEB7369 (<http://www.ebi.ac.uk/ena/data/view/PRJEB7369>).

**Analysis of genomes from isolates.** Filtered and trimmed sequences from the isolated ESBL-producing *Escherichia coli* genomes were assembled using the SPAdes assembler (version 2.5.1) with the additional parameter “-m 32” (6). Resistance genes were identified by matching the assembled

contigs against the Resqu database version 1.1 [www.1928diagnostics.com/resdb](http://www.1928diagnostics.com/resdb) containing 3,019 non-redundant protein sequences corresponding to 325 resistance gene families using blastx (7). See Table S5 for the complete list of genes in Resqu. Only matches longer than 100 bp and with 90% identity or higher were kept. Identified beta-lactam resistance genes were further characterized by BLAST against the entire NCBI protein database, to determine if variants were known or novel.

**rRNA analysis and resistance gene mapping.** Quality filtered read pairs were scanned with Metaxa2 version 2.0 (8) to extract bacterial 16S rRNA (SSU) sequences (additional option “--align none”). In each library, the 16S counts for different genera, families and phyla were normalized to the total number of 16S counts in that library, yielding relative abundances for each organism group. Enterotypes (9) were determined using Principal Component Analysis (PCA) to mainly be driven by the abundances of the *Prevotella* and *Bacteroides* genera, and therefore we defined enterotypes by the ratio of reads mapping to *Prevotella* 16S rRNA sequences and reads mapping to *Bacteroides* 16S rRNA. If the ratio was above one or below one in both the before and after samples from the same individual, the enterotypes was deemed unchanged by travel in that individual. Quality filtered reads were mapped against the Resqu antibiotic resistance gene database using Vmatch (<http://vmatch.de>), allowing two mismatched amino acids per translated read to account for sequencing errors but retain stringency with regards to finding the sought target genes (options “-showdesc 60 -dnvsprot 11 -l 20 -h 2”), and thus avoiding inflation by false positive matches (10). The number of matches to each resistance gene was counted and normalized to the length of the respective gene to avoid bias due to differential length of the resistance genes. The length-normalized numbers were then used for deriving an approximation of the number of resistance genes per 16S rRNA sequence identified by Metaxa2 (see above). The length-normalized numbers were divided by the length of the 16S gene.

***De novo* metagenomic assembly.** *De novo* assembly of the sequences from each sample was performed using Ray Méta (11) on 512 cores of a Cray XE6 system at the PDC Center for High Performance Computing at the Royal Institute of Technology in Stockholm. The k-mer length parameter k was set to 49 based on trials using a range of values in Ray and Velvet (12) assemblies.

**Annotation of assemblies.** Prodigal (13) version 2.60 with parameters “-c -m -p meta” was used to identify open reading frames in assembled contigs and translate those into amino acid sequences. These were subsequently mapped using blastp (in NCBI Blast (7) version 2.2.28+) to the Resque resistance gene database in order to identify contigs containing sequences resembling known resistance genes. Networks of co-localized resistance genes were generated in Cytoscape version 3.0 (14). Contigs were additionally mapped to a reference collection of gastrointestinal microbial genomes to discern taxonomic origin (see below).

**Mapping to reference genomes and contigs.** Quality-filtered reads and assembled contigs were mapped against the Human Microbiome Project (HMP) gastrointestinal\_tract reference genome collection ([http://downloads.hmpdacc.org/data/reference\\_genomes/body\\_sites/Gastrointestinal\\_tract.nuc.fsa](http://downloads.hmpdacc.org/data/reference_genomes/body_sites/Gastrointestinal_tract.nuc.fsa)) using Burrows-Wheeler Aligner, bwa (15), version 0.7.5a with the “mem” subcommand. Reads from all libraries were also mapped back to the total collection of contigs from all samples using Bowtie2 (v. 2.0.2) to estimate their abundance in different libraries.

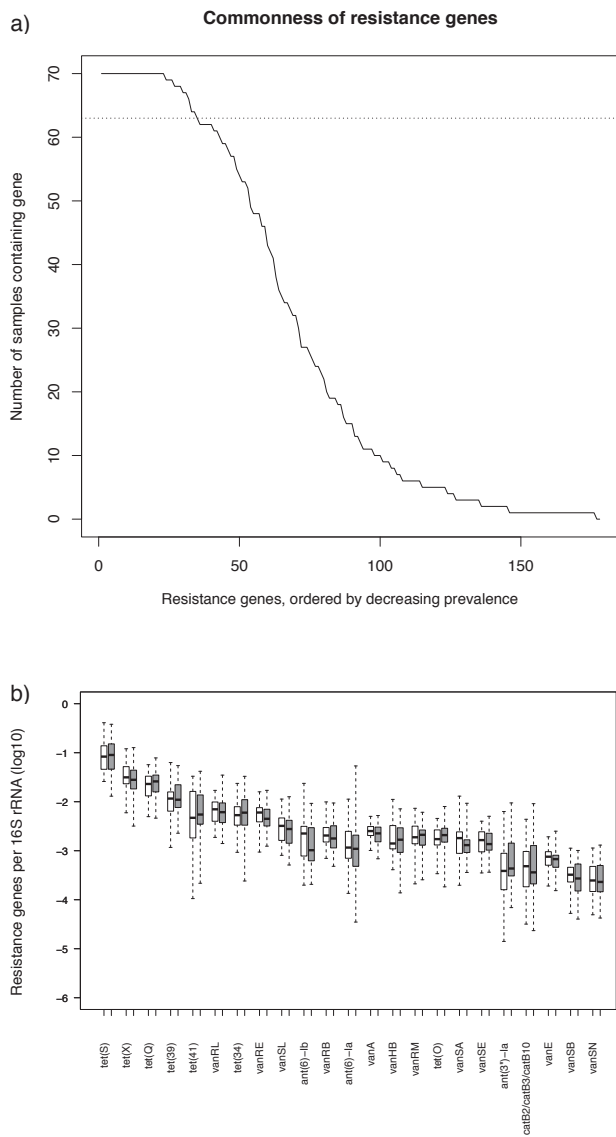
Statistical testing. Resistance gene copies per bacterial 16S rRNA counts were log-transformed by adding a number corresponding to the smallest relative abundance for any detected gene across all samples, and significant changes of average abundances between before and after samples were assessed using paired Student's t-tests. Wilcoxon signed-rank test was used as a complement to find genes with changed median abundance. All p-values were corrected for multiple testing using a Benjamini-Hochberg False Discovery Rate (FDR) and genes with an FDR <0.05 were considered significant (16). The same procedure was adapted for resistance gene categories. Changes in normalized log-transformed relative abundances of taxonomic groups were similarly tested using paired Student's t-tests and tests with an FDR of <0.05 were considered significant. Contigs with significantly different abundance between the groups of travelers returning from the Indian peninsula and Africa were identified using the Limma package (17). Correlations between resistance gene abundances and other factors (gender, age, length of visit, time between return from trip and sample delivery, health care work, sickness during travel, diarrhea, malaria prophylaxis and ESBL culturing results) were assessed using linear regression in the R statistical program (<http://www.r-project.org>). The p-values for each coefficient were corrected for multiple testing and tests with an FDR <0.05 were considered significant.

## References

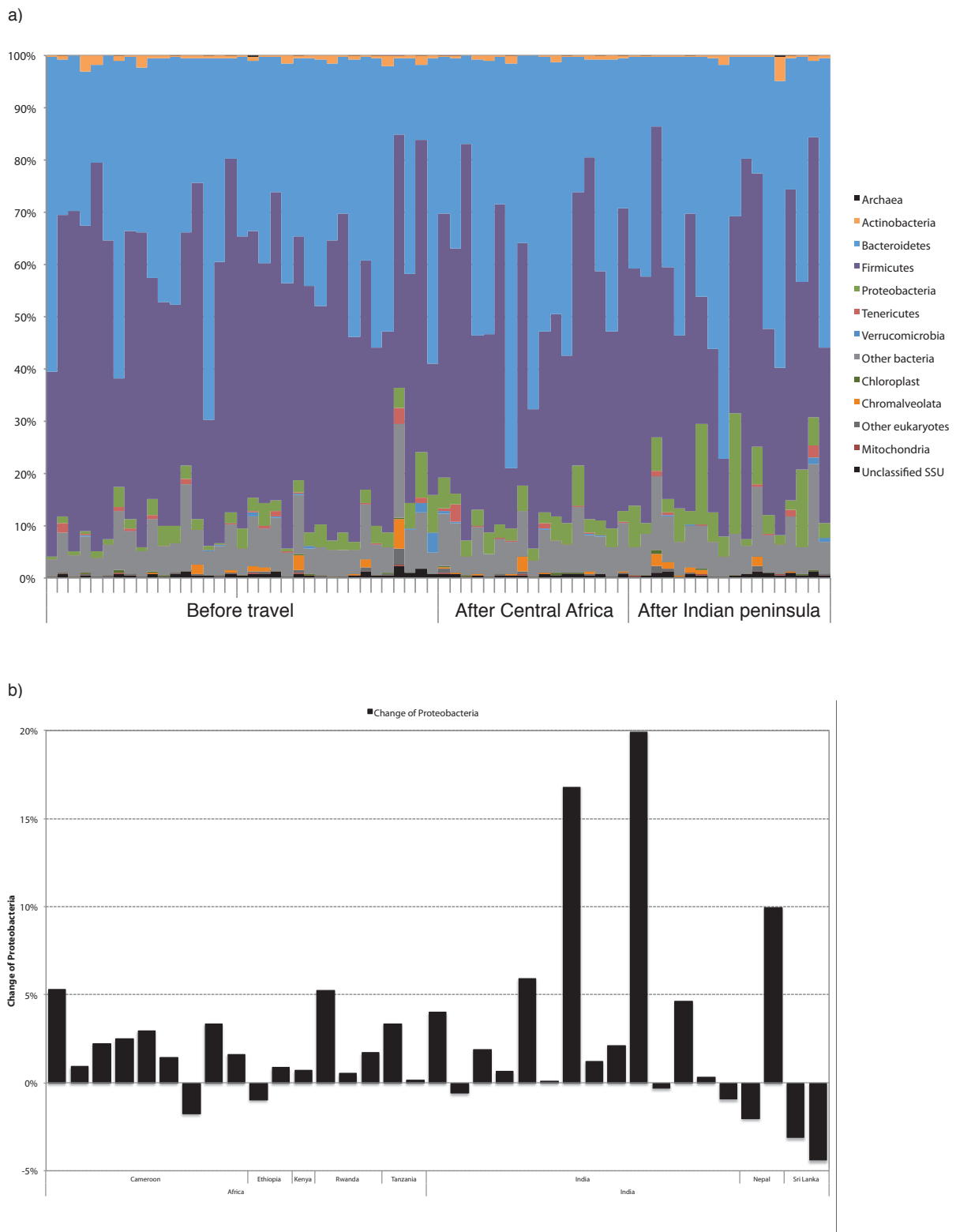
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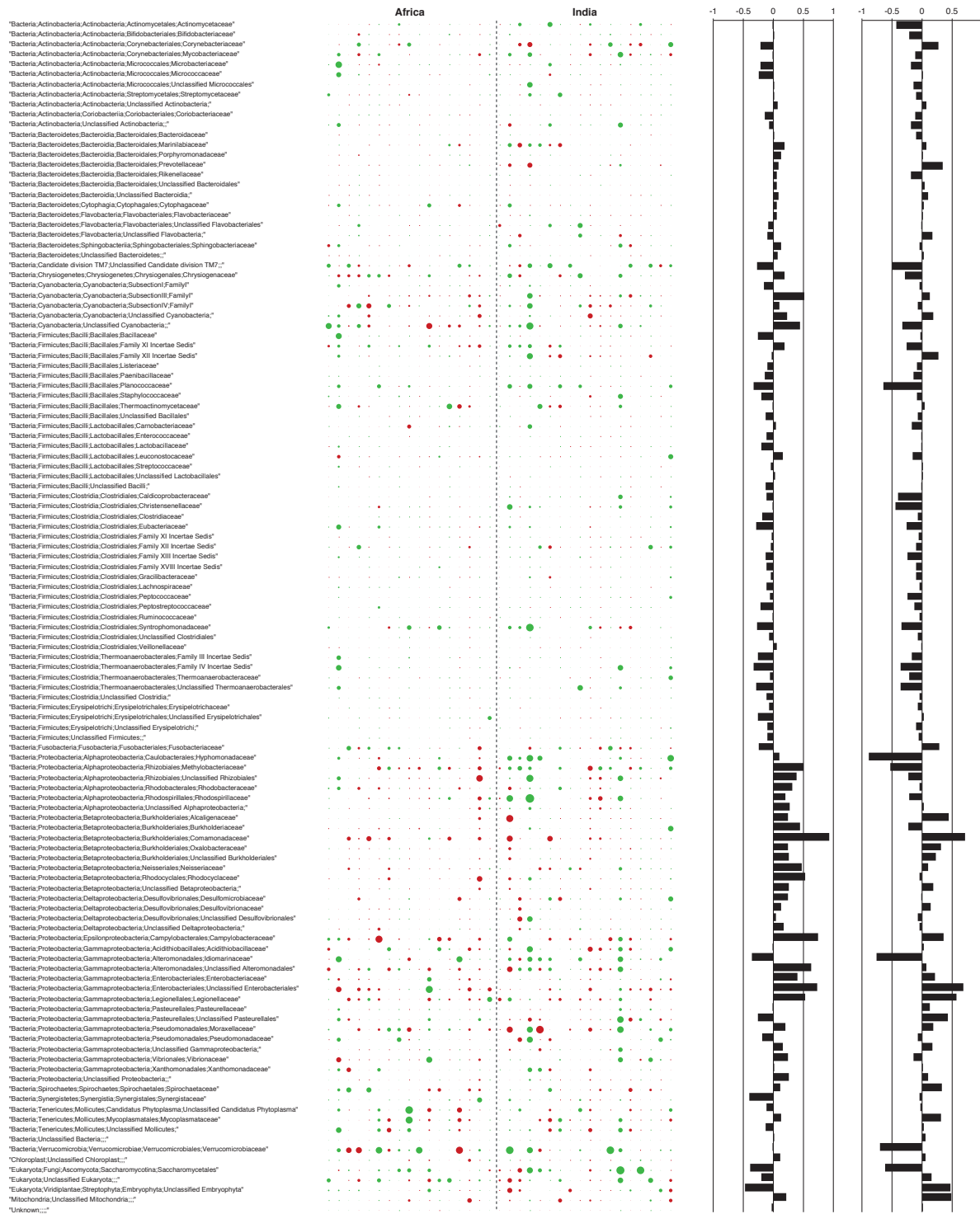
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**Figure S1.** The core resistome of the cohort. a) The number of samples containing a certain resistance gene. b) Abundance of the resistance genes detected in all samples (log 10 scale).





**Figure S3.** Changes of the most abundant taxonomic families, for each individual. The diameter of each dot represents the magnitude of change in that individual (log<sub>10</sub> scale). Green color indicates decreases and red color indicates increases. To the right, the mean fold changes across all individuals traveling to central Africa (left) and the Indian peninsula (right) are shown (log<sub>10</sub> scale).



**Figure S4.** Changes of the most abundant genera, for each individual. The diameter of each dot represents the magnitude of change in that individual (log<sub>10</sub> scale). Green color indicates decreases and red color indicates increases.

Table S1. Study subject and travel characteristics

Region	Destination	Field of study	Days before departure	Length of stay	Days after return	Sex	Age	Patient work in hospital	Travellers diarrhoea	Days ill	Sought medical care	Admitted	Anti-malarial prophylaxis	Mefloquine
Central Africa	Cameroon	Medical	76	49	16	Female	23							x
	Cameroon	Medical	3	29	39	Male	25	x	x	3				x
	Cameroon	Medical	3	30	38	Male	28	x	x	2				x
	Cameroon	Medical	69	49	31	Female	23		x	N/A				x
	Cameroon	Medical	19	34	18	Female	24	x					x	x
	Cameroon	Medical	0	37	35	Female	25		x	3				x
	Cameroon	Medical	41	34	16	Female	25	x	x	6				x
	Cameroon	Medical	25	34	18	Female	28	x	x	4				x
	Cameroon	Medical	11	33	13	Male	25	x	x	1				x
	Ethiopia	Medical	0	150	11	Female	28	x	x	4				x
Ethiopia	Dentistry	16	32	51	Female	24		x	6		x		x	x
Kenya	Medical	3	19	61	Female	28							x	
Tanzania	Medical	0	31	21	Female	24	x			12			x	
Tanzania	Nursing	18	47	32	Male	25	x							x
Rwanda	Medical	2	22	25	Female	25	x		x	15			x	
Rwanda	Medical	2	22	73	Female	25	x		x	1			x	
Rwanda	Medical	2	21	41	Female	26							x	
			Range 0-76 Median 3	Range 19-150 Median 33	Range 11-73 Median 31	76% female (13/17)	Range 23-28 Median 25	65% (11/17)	71% (12/17)	Range 1-15 Median 4	6% (1/17)	0% 0	41% (7/17)	65% (11/17)
Indian peninsula	India	Medical	2	36	5	Male	24	x	x	5				
	India	Medical	15	33	21	Female	26	x					x	
	India	Medical	10	29	19	Female	25	x	x	2			x	
	India	Medical	23	29	44	Female	25	x	x	3				
	India	Medical	4	98	25	Male	23		x	1				
	India	Medical	1	38	2	Female	29	x	x	5			x	
	India	Medical	0	119	22	Female	23	x	x	7			x	
	India	Medical	77	106	20	Female	23	x	x	5				
	India	Medical	10	40	120	Female	22		x	1		x		x
	India	Medical	0	105	19	Female	23		x	2		x		x
India	Medical	16	32	20	Male	24		x	5		x			
India	Medical	1	43	25	Female	27		x	1					
India	Midwife	7	58	32	Female	28		x						
India	Medical	4	53	12	Male	26								
Sri Lanka	Medical	7	23	47	Female	34							x	
Sri Lanka	Medical	9	24	25	Female	24		x						x
Nepal	Medical	4	14	4	Male	28		x						
Nepal	Medical	4	14	4	Female	29		x	x	11				
			Range 0-77 Median 6	Range 14-119 Median 37	Range 2-120 Median 21	72% female (13/18)	Range 23-34 Median 25	67% (12/18)	67% (12/18)	Range 1-7 Median 4	11% (2/18)	6% (1/18)	28% (5/18)	22% (4/18)

Table S2. Details of the individual sequencing libraries

Location data		SSU sequences identified by Metaxa2										Library sizes			
Before or after	Destination	Archaea	Bacteria	Chloroplast	Eukaryota	Mitochondria	Unknown	Raw library size (reads)	Raw library size (Gbp)	Quality-filtered size	Low quality reads	Bacterial 16S per million reads			
Before	Cameroon	0	220,291	22	157	9	433	125,907,692	25.18	116,985,641	8,922,051	1,883.06			
Before	Cameroon	23	121,442	47	141	97	929	78,430,878	15.69	70,967,328	7,463,550	1,711.24			
Before	Cameroon	0	182,229	16	24	23	506	110,624,462	22.12	102,862,162	7,762,300	1,771.58			
Before	Cameroon	0	203,600	157	703	38	1,010	138,278,024	27.66	129,038,382	9,239,642	1,577.83			
Before	Cameroon	0	189,107	40	97	20	265	97,108,136	19.42	84,768,485	12,339,651	2,230.86			
Before	Cameroon	0	208,475	231	394	45	585	105,488,039	21.10	96,338,305	9,149,734	2,163.99			
Before	Cameroon	15	217,277	1,113	515	38	1,947	94,583,689	18.92	86,709,535	7,874,154	2,505.80			
Before	Cameroon	0	112,218	8	217	4	578	57,708,298	11.54	55,274,739	2,433,559	2,030.19			
Before	Cameroon	0	199,666	54	141	20	429	93,610,789	18.72	80,289,041	13,321,748	2,486.84			
Before	Ethiopia	0	198,146	394	513	22	1,480	114,623,643	22.92	107,097,525	7,526,118	1,850.15			
Before	Ethiopia	0	208,188	355	1	2	576	96,158,167	19.23	94,094,185	2,063,982	2,212.55			
Before	Kenya	17	222,448	44	71	36	1,858	152,444,626	30.49	143,428,931	9,015,695	1,590.93			
Before	Tanzania	0	184,030	19	433	4	596	94,496,474	18.90	91,204,324	3,292,150	2,017.78			
Before	Tanzania	84	363,035	88	2,734	80	2,848	186,020,932	37.20	178,004,607	8,016,325	2,039.47			
Before	Rwanda	0	136,406	177	9	1	1,635	79,870,933	15.97	73,992,049	5,878,884	1,843.52			
Before	Rwanda	0	111,466	21	2,103	184	496	58,683,097	11.74	56,688,930	1,994,167	1,966.27			
Before	Rwanda	2	248,514	1	12	0	1,140	116,336,923	23.27	112,218,587	4,118,336	2,214.55			
Before	India	0	164,203	74	100	16	1,907	94,168,067	18.83	86,725,925	7,442,142	1,893.36			
Before	India	0	213,993	83	6	1	530	125,452,729	25.09	120,493,836	4,958,893	1,775.97			
Before	India	9	123,055	166	4,387	151	1,116	81,137,887	16.23	77,796,173	3,341,714	1,581.76			
Before	India	0	219,475	6	7	4	529	105,390,115	21.08	99,898,565	5,491,550	2,196.98			
Before	India	12	108,377	1	192	7	487	68,826,005	13.77	65,736,512	3,089,493	1,648.66			
Before	India	0	187,490	159	4,177	97	2,485	126,672,111	25.33	117,356,765	9,315,346	1,597.61			
Before	India	16	215,319	56	197	24	1,051	114,873,991	22.97	110,841,646	4,032,345	1,942.58			
Before	India	0	191,903	37	157	41	744	108,862,964	21.77	100,505,030	8,357,934	1,909.39			
Before	India	224	152,457	140	2,199	143	1,277	89,041,296	17.81	84,176,188	4,865,108	1,811.17			
Before	India	0	180,204	335	2,298	72	1,200	104,860,302	20.97	97,572,943	7,287,359	1,846.86			
Before	India	0	144,228	36	269	68	554	119,776,667	23.96	111,456,534	8,320,133	1,294.03			
Before	India	0	154,785	32	334	67	410	108,484,366	21.70	102,117,614	6,366,752	1,515.75			
Before	India	0	164,743	8	29	13	499	108,191,129	21.64	98,753,669	9,437,460	1,668.22			
Before	India	1	200,324	114	821	52	913	120,944,486	24.19	111,299,634	9,644,852	1,799.86			
Before	Sri Lanka	146	137,803	214	13	4	2,439	101,623,663	20.32	97,424,636	4,199,027	1,414.46			
Before	Sri Lanka	0	196,377	24	109	0	1,423	180,821,242	36.16	168,578,941	12,242,301	1,164.90			
Before	Nepal	11	100,230	198	9,846	251	2,677	81,153,547	16.23	75,267,243	5,886,304	1,331.65			
Before	Nepal	0	141,454	18	8	0	1,292	97,563,912	19.51	91,026,911	6,537,001	1,553.98			
After	Cameroon	5	112,023	169	1,382	239	857	68,589,271	13.72	64,867,085	3,722,186	1,726.96			
After	Cameroon	2	160,097	70	275	32	1,169	92,482,953	18.50	85,135,965	7,346,988	1,880.49			
After	Cameroon	0	149,056	21	15	12	513	95,181,317	19.04	80,837,880	14,343,437	1,843.89			

After	Cameroon	0	182,566	71	719	46	760	108,827,882	21.77	94,736,141	14,091,741	1,927.10
After	Cameroon	0	157,990	10	5	2	310	112,558,957	22.51	105,987,973	6,570,984	1,490.64
After	Cameroon	0	185,416	27	557	62	914	113,933,461	22.79	105,653,830	8,279,631	1,754.94
After	Cameroon	6	215,167	312	376	24	1,047	74,880,136	14.98	71,943,878	2,936,258	2,990.76
After	Cameroon	0	206,543	372	6,979	630	1,179	127,104,270	25.42	119,741,546	7,362,724	1,724.91
After	Cameroon	0	264,177	7	24	15	558	138,551,617	27.71	124,497,835	14,053,782	2,121.94
After	Ethiopia	0	258,280	45	218	20	2,127	148,483,688	29.70	139,510,543	8,973,145	1,851.33
After	Ethiopia	0	54,397	294	42	2	287	82,739,446	16.55	40,538,343	42,201,103	1,341.87
After	Kenya	20	168,496	10	191	18	1,348	118,039,879	23.61	110,534,095	7,505,784	1,524.38
After	Tanzania	0	264,522	59	86	80	690	132,336,558	26.45	127,715,066	4,521,492	2,071.19
After	Tanzania	8	100,527	160	420	30	798	463,791,736	92.76	54,987,641	408,804,095	1,828.17
After	Rwanda	0	243,606	352	1	2	1,885	137,856,261	27.57	127,235,892	10,620,369	1,914.60
After	Rwanda	0	204,169	31	1,718	243	865	98,323,380	19.66	92,301,898	6,021,482	2,211.97
After	Rwanda	14	169,981	102	111	20	1,267	108,508,604	21.70	103,146,520	5,362,084	1,647.96
After	India	0	308,378	152	5,518	131	3,696	172,298,973	34.46	164,171,030	8,127,943	1,878.39
After	India	0	213,865	46	244	30	600	95,916,677	19.18	92,069,410	3,847,267	2,322.87
After	India	1	271,377	35	3,645	226	1,820	145,856,676	29.17	139,769,906	6,086,770	1,941.60
After	India	0	156,408	4	5	7	598	80,858,580	16.17	77,801,453	3,057,127	2,010.35
After	India	0	186,891	13	79	7	1,271	103,809,366	20.76	95,299,798	8,509,568	1,961.08
After	India	0	196,638	143	5,629	140	2,428	119,333,838	23.87	111,437,652	7,896,186	1,764.56
After	India	0	142,048	37	26	6	1,494	79,961,086	15.99	74,749,858	5,211,228	1,900.31
After	India	0	191,360	16	324	277	585	114,439,546	22.89	80,387,488	34,052,058	2,380.47
After	India	6	123,942	42	66	47	621	78,223,579	15.64	75,053,057	3,170,522	1,651.39
After	India	0	208,535	1,555	8,091	178	2,001	103,437,332	20.69	94,520,827	8,916,505	2,206.23
After	India	0	228,654	191	2,104	466	1,308	109,989,613	22.00	102,842,328	7,147,285	2,223.35
After	India	0	131,170	6	1	10	158	70,806,355	14.16	68,666,720	2,139,635	1,910.24
After	India	0	96,110	2	28	3	91	77,982,751	15.60	75,870,339	2,112,412	1,266.77
After	India	197	137,542	47	105	81	838	101,742,471	20.35	89,404,424	12,338,047	1,538.42
After	Sri Lanka	35	163,143	301	5	6	2,155	101,637,724	20.33	97,729,754	3,907,970	1,669.33
After	Sri Lanka	0	108,658	53	32	7	663	85,438,843	17.09	81,634,162	3,804,681	1,331.04
After	Nepal	35	221,127	157	137	24	2,394	136,599,459	27.32	125,704,413	10,895,046	1,759.10
After	Nepal	0	93,541	22	21	3	567	58,982,996	11.80	51,705,957	7,277,039	1,809.10





**Table S4.** Beta-lactam resistance genes found in the genomes of ESBL-producing *E. coli*

<b>Individual</b>	<b>CTX-M</b>	<b>OXA-1</b>	<b>SHV</b>	<b>TEM</b>
<b>103</b>	x	x	x	
<b>107</b>	x	x		x
<b>120</b>	x	x		
<b>139</b>	x	x		x
<b>147</b>	x			x
<b>154</b>	x	x		x
<b>155</b>	x			
<b>170</b>	x			x
<b>176</b>	x	x	x	x
<b>177</b>	x			x
<b>183</b>	x			x
<b>184</b>	x	x		x
<b>211<sup>a</sup></b>	x	x		x

<sup>a</sup> Individual 211 had ESBL-producing *E. coli* before travel, but not after.

**Table S5.** Resistance genes included in the Resqu database

<b>Gene family</b>	<b>Classification</b>	<b>Resistance</b>
ACC	Ambler class C	Beta-lactam
CTX-M	Ambler class A	Beta-lactam
CAR	Ambler class A	Beta-lactam
OXA-63	Ambler class D	Beta-lactam
OXA-62	Ambler class D	Beta-lactam
PER	Ambler class A	Beta-lactam
OXA-60	Ambler class D	Beta-lactam
ACT	Ambler class C	Beta-lactam
SME	Ambler class A	Beta-lactam
MIR	Ambler class C	Beta-lactam
GES	Ambler class A	Beta-lactam
TEM	Ambler class A	Beta-lactam
CMY1	Ambler class C	Beta-lactam
OXA-55	Ambler class D	Beta-lactam
NDM	Ambler class B	Beta-lactam
OXA-50	Ambler class D	Beta-lactam
OXA-51	Ambler class D	Beta-lactam
CMY2	Ambler class C	Beta-lactam
OXA-10	Ambler class D	Beta-lactam
FOX	Ambler class C	Beta-lactam
OXA-58	Ambler class D	Beta-lactam
OXA-48	Ambler class D	Beta-lactam
SHV	Ambler class A	Beta-lactam
OXA-1	Ambler class D	Beta-lactam
OXA-2	Ambler class D	Beta-lactam
KPC	Ambler class A	Beta-lactam
VEB	Ambler class A	Beta-lactam
DHA	Ambler class C	Beta-lactam
IND	Ambler class B	Beta-lactam
MOX	Ambler class C	Beta-lactam
IMP	Ambler class B	Beta-lactam
CFE	Ambler class C	Beta-lactam
CAR (RTG)	Ambler class A	Beta-lactam
VIM	Ambler class B	Beta-lactam
OXA-23	Ambler class D	Beta-lactam
OXA-20	Ambler class D	Beta-lactam
OXA-24	Ambler class D	Beta-lactam
LAT	Ambler class C	Beta-lactam
aac(6')-Ig	acetyltransferase	Aminoglycoside
aac(6')-If	acetyltransferase	Aminoglycoside
aac(6')-Ia	acetyltransferase	Aminoglycoside
aac(6')-Ic	acetyltransferase	Aminoglycoside
aac(6')-Ib	acetyltransferase	Aminoglycoside
aac(6')-Im	acetyltransferase	Aminoglycoside
aac(6')-Il	acetyltransferase	Aminoglycoside
aph(6)-Ia	phosphotransferase	Aminoglycoside
ant(6)-Ib	nucleotidyltransferase	Aminoglycoside
aac(6')-Ii	acetyltransferase	Aminoglycoside
aph(6)-Ib	phosphotransferase	Aminoglycoside

aac(6')-Ik	acetyltransferase	Aminoglycoside
aac(6')-Ij	acetyltransferase	Aminoglycoside
ant(9)-Ib	nucleotidyltransferase	Aminoglycoside
ant(6)-Ia	nucleotidyltransferase	Aminoglycoside
aac(6')-Ir	acetyltransferase	Aminoglycoside
aac(6')-IIa	acetyltransferase	Aminoglycoside
aac(6')-29a/29b	acetyltransferase	Aminoglycoside
aac(6')-IIc	acetyltransferase	Aminoglycoside
aac(3)-Ie	acetyltransferase	Aminoglycoside
aac(3)-Ib	acetyltransferase	Aminoglycoside
aac(3)-Ic	acetyltransferase	Aminoglycoside
aac(3)-Ia	acetyltransferase	Aminoglycoside
aac(2')-Ia	acetyltransferase	Aminoglycoside
ant(3'')-Ia	nucleotidyltransferase	Aminoglycoside
aac(2')-Ic	acetyltransferase	Aminoglycoside
aac(6')-IIb	acetyltransferase	Aminoglycoside
aph(3')-Vc	phosphotransferase	Aminoglycoside
aac(6')-32	acetyltransferase	Aminoglycoside
aac(3)-VIa	acetyltransferase	Aminoglycoside
aph(3')-Vb	phosphotransferase	Aminoglycoside
aac(6')-Iai	acetyltransferase	Aminoglycoside
aph(3')-VIIa	phosphotransferase	Aminoglycoside
aac(3)-IXa	acetyltransferase	Aminoglycoside
aph(4)-Ia	phosphotransferase	Aminoglycoside
aph(3')-VIa	phosphotransferase	Aminoglycoside
aph(4)-Ib	phosphotransferase	Aminoglycoside
aac(3)-IIIb/IIIc	acetyltransferase	Aminoglycoside
aph(3'')-Ic	phosphotransferase	Aminoglycoside
aac(6')-sk	acetyltransferase	Aminoglycoside
ant(4')-Ia	nucleotidyltransferase	Aminoglycoside
aph(6)-Id	phosphotransferase	Aminoglycoside
aac(6')-Iae	acetyltransferase	Aminoglycoside
aac(6')-Iad	acetyltransferase	Aminoglycoside
aph(3')-IIa	phosphotransferase	Aminoglycoside
aac(3)-IVa	acetyltransferase	Aminoglycoside
aph(6)-Ic	phosphotransferase	Aminoglycoside
aph(3')-Va	phosphotransferase	Aminoglycoside
ant(4')-IIa	nucleotidyltransferase	Aminoglycoside
aph(3')-IIb	phosphotransferase	Aminoglycoside
ant(4')-IIb	nucleotidyltransferase	Aminoglycoside
aph(9)-Ia	phosphotransferase	Aminoglycoside
aac(3)-IIb	acetyltransferase	Aminoglycoside
aac(2')-Ib	acetyltransferase	Aminoglycoside
aac(6')-Iy/z/aa	acetyltransferase	Aminoglycoside
aph(3'')-Ib	phosphotransferase	Aminoglycoside
aph(3')-IVa	phosphotransferase	Aminoglycoside
aac(3)-IIa/IIc	acetyltransferase	Aminoglycoside
aac(2')-Id	acetyltransferase	Aminoglycoside
aac(3)-VIIIa	acetyltransferase	Aminoglycoside
aac(6')-Ip/q/af	acetyltransferase	Aminoglycoside
aac(3)-VIIa	acetyltransferase	Aminoglycoside
aph(3'')-Ia	phosphotransferase	Aminoglycoside

ant(9)-Ia	nucleotidyltransferase	Aminoglycoside
aph(3')-IIIa	phosphotransferase	Aminoglycoside
aac(3)-Xa	acetyltransferase	Aminoglycoside
aph(9)-Ib	phosphotransferase	Aminoglycoside
aac(6')-Iid	acetyltransferase	Aminoglycoside
aac(6')-Iih	acetyltransferase	Aminoglycoside
aph(2'')-Ie/IVa	phosphotransferase	Aminoglycoside
aac(3)-IIIa	acetyltransferase	Aminoglycoside
aph(2'')-IIIa	phosphotransferase	Aminoglycoside
aph(3')-Ib	phosphotransferase	Aminoglycoside
aph(2'')-IIa	phosphotransferase	Aminoglycoside
aph(3')-Ia	phosphotransferase	Aminoglycoside
aac(6')-33/130	acetyltransferase	Aminoglycoside
aac(6')-Ih/Is/Iu/Iv/Iw/Ix	acetyltransferase	Aminoglycoside
ant(2'')-Ia	nucleotidyltransferase	Aminoglycoside
qnrC	pentapeptide repeat protein class C	Quinolone
qnrB	pentapeptide repeat protein class B	Quinolone
qnrA	pentapeptide repeat protein class A	Quinolone
qnrS	pentapeptide repeat protein class S	Quinolone
qnrD	pentapeptide repeat protein class D	Quinolone
smr/qac/abr	efflux pump	Multidrug
qacAB	efflux pump	Multidrug
nhsA	methyltransferase	Thiostrepton
tsnRA	methyltransferase	Thiostrepton
tsnRB	methyltransferase	Thiostrepton
catA1	acetyltransferase	Amphenicol
catA2	acetyltransferase	Amphenicol
catA3	acetyltransferase	Amphenicol
catA4	acetyltransferase	Amphenicol
catA5	acetyltransferase	Amphenicol
catA6	acetyltransferase	Amphenicol
catA7/catA8	acetyltransferase	Amphenicol
catA9	acetyltransferase	Amphenicol
catA10	acetyltransferase	Amphenicol
catA11	acetyltransferase	Amphenicol
catA12	acetyltransferase	Amphenicol
catA13	acetyltransferase	Amphenicol
catA15	acetyltransferase	Amphenicol
catA16	acetyltransferase	Amphenicol
catB1	acetyltransferase	Amphenicol
catB2/catB3/catB10	acetyltransferase	Amphenicol
catB7	acetyltransferase	Amphenicol
catB9	acetyltransferase	Amphenicol
cmlA	efflux pump	Amphenicol
cml-II	efflux pump	Amphenicol
floR	efflux pump	Amphenicol
fexA	efflux pump	Amphenicol
cml-III	efflux pump	Amphenicol
cmlv	efflux pump	Amphenicol
cmrA	efflux pump	Amphenicol
cmr-II	efflux pump	Amphenicol
pexA	efflux pump	Amphenicol

erm(A)	methylase	Macrolide
erm(B)	methylase	Macrolide
erm(C)	methylase	Macrolide
erm(D)	methylase	Macrolide
erm(E)	methylase	Macrolide
erm(F)	methylase	Macrolide
erm(G)	methylase	Macrolide
erm(H)	methylase	Macrolide
erm(N)	methylase	Macrolide
erm(O)	methylase	Macrolide
erm(Q)	methylase	Macrolide
erm(R)	methylase	Macrolide
erm(S)	methylase	Macrolide
erm(T)	methylase	Macrolide
erm(U)	methylase	Macrolide
erm(V)	methylase	Macrolide
erm(W)	methylase	Macrolide
erm(X)	methylase	Macrolide
erm(Y)	methylase	Macrolide
erm(Z)	methylase	Macrolide
erm(30)	methylase	Macrolide
erm(31)	methylase	Macrolide
erm(32)	methylase	Macrolide
erm(33)	methylase	Macrolide
erm(34)	methylase	Macrolide
erm(35)	methylase	Macrolide
erm(36)	methylase	Macrolide
erm(41)	methylase	Macrolide
erm(42)	methylase	Macrolide
cfr	methylase	Macrolide
msr(A)	efflux pump	Macrolide
vga(A)	efflux pump	Macrolide
vga(B)	efflux pump	Macrolide
vga(C)	efflux pump	Macrolide
ere(A)/ere(C)	esterase	Macrolide
ere(B)	esterase	Macrolide
vgb(A)	lyase	Macrolide
vgb(B)	lyase	Macrolide
lnu(A)	transferase	Macrolide
lnu(B)	transferase	Macrolide
lnu(C)	transferase	Macrolide
lnu(D)	transferase	Macrolide
lnu(F)	transferase	Macrolide
vat(A)	transferase	Macrolide
vat(B)	transferase	Macrolide
vat(C)	transferase	Macrolide
vat(D)	transferase	Macrolide
vat(E)	transferase	Macrolide
vat(F)	transferase	Macrolide
vat(G)	transferase	Macrolide
mph(A)	phosphorylase	Macrolide
mph(B)	phosphorylase	Macrolide

mph(C)	phosphorylase	Macrolide
mph(E)	phosphorylase	Macrolide
cmr	NA	Macrolide
mdt(A)	efflux pump	Macrolide
qepA	efflux pump	quinolone
sul1	dihydropteroate synthetase inhibitor	sulfonamide
sul2	dihydropteroate synthetase inhibitor	sulfonamide
sul3	dihydropteroate synthetase inhibitor	sulfonamide
otr(A)		Tetracycline
tetB(P)		Tetracycline
tet(M)		Tetracycline
tet(O)		Tetracycline
tet(Q)		Tetracycline
tet(S)		Tetracycline
tet(T)		Tetracycline
tet(W)		Tetracycline
tet(32)		Tetracycline
tet(36)		Tetracycline
tet(44)		Tetracycline
otr(B)	efflux pump	Tetracycline
otr(C)	efflux pump	Tetracycline
tcr	efflux pump	Tetracycline
tet(A)	efflux pump	Tetracycline
tetA(P)	efflux pump	Tetracycline
tet(B)	efflux pump	Tetracycline
tet(C)	efflux pump	Tetracycline
tet(D)	efflux pump	Tetracycline
tet(E)	efflux pump	Tetracycline
tet(G)	efflux pump	Tetracycline
tet(H)	efflux pump	Tetracycline
tet(J)	efflux pump	Tetracycline
tet(K)	efflux pump	Tetracycline
tet(L)	efflux pump	Tetracycline
tet(V)	efflux pump	Tetracycline
tet(Y)	efflux pump	Tetracycline
tet(Z)	efflux pump	Tetracycline
tet(30)	efflux pump	Tetracycline
tet(31)	efflux pump	Tetracycline
tet(33)	efflux pump	Tetracycline
tet(39)	efflux pump	Tetracycline
tet(40)	efflux pump	Tetracycline
tet(41)	efflux pump	Tetracycline
tet(42)	efflux pump	Tetracycline
tet(43)	efflux pump	Tetracycline
tet(X)		Tetracycline
tet(34)		Tetracycline
tet(37)		Tetracycline
tet(U)		Tetracycline
dfrA1/dfrA15	dihydrofolate reductase	Sufonamide
dfrA3	dihydrofolate reductase	Sufonamide
dfrA5/dfrA14/dfrA25/dfrA30	dihydrofolate reductase	Sufonamide
dfrA6/dfrA31	dihydrofolate reductase	Sufonamide

dfrA7/dfrA17/dfrA32	dihydrofolate reductase	Sulfonamide
dfrA8	dihydrofolate reductase	Sulfonamide
dfrA9	dihydrofolate reductase	Sulfonamide
dfrA10	dihydrofolate reductase	Sulfonamide
dfrA12/dfrA13/dfrA21/dfrA22/dfrA33	dihydrofolate reductase	Sulfonamide
dfrA16	dihydrofolate reductase	Sulfonamide
dfrA18	dihydrofolate reductase	Sulfonamide
dfrA20	dihydrofolate reductase	Sulfonamide
dfrA23	dihydrofolate reductase	Sulfonamide
dfrA24	dihydrofolate reductase	Sulfonamide
dfrA26	dihydrofolate reductase	Sulfonamide
dfrA27/dfrA28	dihydrofolate reductase	Sulfonamide
dfrB1/dfrB5/dfrB6/dfrB8	dihydrofolate reductase	Sulfonamide
dfrB2/dfrB3/dfrB7	dihydrofolate reductase	Sulfonamide
dfrB4	dihydrofolate reductase	Sulfonamide
dfrC	dihydrofolate reductase	Sulfonamide
dfrD	dihydrofolate reductase	Sulfonamide
dfrG	dihydrofolate reductase	Sulfonamide
dfrK	dihydrofolate reductase	Sulfonamide
vanA	ligase of type A	Glycopeptide
vanRA	transcriptional regulator of type A	Glycopeptide
vanSA	histidine protein kinase of type A	Glycopeptide
vanHA	dehydrogenase of type A	Glycopeptide
vanXA	dipeptidase of type A	Glycopeptide
vanYA	carboxypeptidase of type A	Glycopeptide
vanZA	teicoplanin resistance protein of type A	Glycopeptide
vanB	ligase of type B	Glycopeptide
vanXB	dipeptidase of type B	Glycopeptide
vanHB	dehydrogenase of type B	Glycopeptide
vanWB	glycopeptide resistance gene of type B	Glycopeptide
vanYB	carboxypeptidase of type B	Glycopeptide
vanSB	transcriptional regulator of type B	Glycopeptide
vanRB	histidine protein kinase of type B	Glycopeptide
vanD	ligase of type D	Glycopeptide
vanXD	dipeptidase of type D	Glycopeptide
vanHD	dehydrogenase of type D	Glycopeptide
vanYD	carboxypeptidase of type D	Glycopeptide
vanSD	transcriptional regulator of type D	Glycopeptide
vanRD	histidine protein kinase of type D	Glycopeptide
vanG	ligase of type D	Glycopeptide
vanXYG	dipeptidase/carboxypeptidase of type G	Glycopeptide
vanWG	glycopeptide resistance gene of type G	Glycopeptide
vanYG	carboxypeptidase of type G	Glycopeptide
vanSG	transcriptional regulator of type G	Glycopeptide
vanRG	histidine protein kinase of type G	Glycopeptide
vanTG	serine racemase of type G	Glycopeptide
vanE	ligase of type E	Glycopeptide
vanXYE	dipeptidase/carboxypeptidase of type E	Glycopeptide
vanTE	serine racemase of type E	Glycopeptide
vanRE	transcriptional regulator of type E	Glycopeptide
vanSE	histidine protein kinase of type E	Glycopeptide
vanL	ligase of type type L	Glycopeptide

vanXYL	dipeptidase/carboxypeptidase of type L	Glycopeptide
vanTmL	serine racemase of type L	Glycopeptide
vanTrL	serine racemase of type L	Glycopeptide
vanRL	transcriptional regulator of type L	Glycopeptide
vanSL	histidine protein kinase of type L	Glycopeptide
vanM	ligase of type M	Glycopeptide
vanXM	dipeptidase of type M	Glycopeptide
vanHM	dehydrogenase of type M	Glycopeptide
vanYM	carboxypeptidase of type M	Glycopeptide
vanSM	histidine protein kinase of type M	Glycopeptide
vanRM	transcriptional regulator of type M	Glycopeptide
vanN	ligase of type N	Glycopeptide
vanXYN	dipeptidase/carboxypeptidase of type N	Glycopeptide
vanTN	serine racemase of type N	Glycopeptide
vanRN	transcriptional regulator of type N	Glycopeptide
vanSN	histidine protein kinase of type N	Glycopeptide
intI9	Integron-associated integrase (class 9)	
intI8	Integron-associated integrase (class 8)	
intI7	Integron-associated integrase (class 7)	
intI6	Integron-associated integrase (class 6)	
intI1	Integron-associated integrase (class 1)	
intI3	Integron-associated integrase (class 3)	
intI2	Integron-associated integrase (class 2)	
intI10	Integron-associated integrase (class 10)	
ISCR8	ISCR transposase	
ISCR3	ISCR transposase	
ISCR4	ISCR transposase	
ISCR5	ISCR transposase	
ISCR14	ISCR transposase	
ISCR7	ISCR transposase	
ISCR1	ISCR transposase	
ISCR2	ISCR transposase	
ISCR6	ISCR transposase	

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