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**Distribution of mutation frequencies among *Salmonella enterica* isolates
from animal and human sources and genetic characterization
of a *Salmonella* Heidelberg hypermutator**

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Abstract

Hypermutation is an important mechanism used by different *Salmonella enterica* subspecies *enterica* to regulate genetic stability in adaptation to changing environments, including antimicrobial treatments and industrial processes. Strong hypermutator strains generally contain a mutation in genes of the methyl mismatch repair (MMR) system and have mutation frequencies up to 1000 fold higher than wild type strains. The objectives of this study were to determine the distribution of mutation frequencies from a collection of 209 *Salmonella* strains, to genetically characterize a strong mutator, and to study MMR mutated protein-DNA binding interactions. Only one strain of *S. Heidelberg* was determined to have a hypermutator phenotype by virtue of its high mutation rate. Sequencing of genes of the MMR system showed a 12 bp deletion in the *mutS* gene was present. The MMR mutated protein-DNA binding interactions were studied by bioanalysis, using the available crystal structure of a similar MutS protein from *Escherichia coli*. This analysis showed the small deletion in the *Salmonella* MutS was localized within the core domain. A retardation assay with MutS from hypermutable and wild type strains showed this mutation has no effect on MutS DNA binding. A better understanding of the genetic mechanisms of hypermutation will help to anticipate the behavior of hypermutator strains in various conditions.

Key words: *Salmonella* Heidelberg; MutS; hypermutation; DNA retardation assay; His6-tag protein.

1. Introduction

Salmonella enterica subspecies *enterica* (*S.*) are transmissible from animals to humans (Vaillant et al., 2005) and many serovars, including *S.* Heidelberg, are responsible for food-related illnesses causing localized gastroenteritis, with increasing rates of resistance to multiple antimicrobial agents (Zhao et al., 2008). Hypermutable bacteria have an elevated mutation rate, up to 1000 fold higher than normal and the frequency of transduction can be 10-15 fold higher than normal (Leclerc et al 1996, Merino et al., 2002), compared to their wild-type counterparts (spontaneous mutator phenotype) (Matic et al., 1997). Hypermutable bacteria play a role in adapting to environmental stresses and thus, in developing antibiotic resistance (Taddei et al., 1997).

The majority of natural bacterial strains with the hypermutator phenotype harbor mutations in the methyl mismatch repair (MMR) system, predominantly in the *mutS* gene (Boe et al., 2000; Chopra et al., 2003; Matic et al., 2006). The essential role of this system is to correct errors that have escaped the proofreading function of polymerase, such as mismatches between base-pairs (bp) or errors due to insertions or deletions (Yang, 2000). This system relies on the activity of four major proteins: MutS, MutL, MutH, and UvrD. MutS proteins are responsible for repairing base mismatches and insertions and deletions up to four bases. MutS proteins also play a role in preventing recombination between non-identical DNA sequences (Radman et al., 1995). In the presence of both ATP and a mismatch, MutS recruits MutL. Together they activate MutH and initiate the entire MMR system. Hypermutable strains have been detected in clinical settings among bacterial strains that colonized the lung of a cystic fibrosis patient (Oliver et al., 2000; Prunier et al., 2003) and the urinary tract (Denamur et al. 2002).

The objectives of this study were (1) to determine the distribution of mutation frequencies from a collection of 209 *Salmonella* strains, of animal and human origin, to select a strain with a hypermutator phenotype by virtue of its high mutation rate; (2) to genetically characterize a strong mutator; and (3) to study MMR mutated protein-DNA binding interactions, using a gel shift assay and a 3D model of *E. coli* MutS protein.

2. Material and methods

2.1. Bacterial strains and phenotypic identification

A total of 209 *Salmonella enterica* strains were isolated from humans (62 clinical isolates from Rennes Hospital, France) and animals (112 cows, 25 pigs, 3 sheep, and 7 chickens, from the Veterinary Laboratory of Ille et Vilaine, France) from 1995 to 2005 (Table 1). Identification was performed with API (API 20E, API 32GN, BioMérieux, Marcy l'Etoile, France). *Escherichia coli* M15 pREP 4 (Qiagen, Courtaboeuf, France) was used for cloning and constructing the His₆-tagged plasmid, as described below.

2.2. Mutation rates

Mutation rates were determined as described previously (LeClerc et al., 1996). Briefly, a single colony was suspended in 10 mL of Luria Bertani (LB; AES Laboratory, Combourg, France) broth and incubated at 37°C for 24 h. This pre-culture was inoculated at a dilution of 1:50 in 5 mL of fresh LB broth and then incubated at 37°C until OD₆₀₀ = 0.6 was reached. One hundred microliters of culture were spread onto LB agar with or without rifampicin (100µg/mL, Sigma Aldrich, Saint Quentin Fallavier, France). To confirm the mutator phenotype, the mutation frequencies that conferred resistance to the antibiotics nalidixic acid (50µg/mL, Sigma Aldrich, Saint Quentin Fallavier, France) and fosfomycin (100µg/mL,

Sigma Aldrich, Saint Quentin Fallavier, France) were determined. Mutation frequencies are reported as a proportion of rifampicin-resistant colonies to the total viable count. The results were related to the mean value obtained from three independent cultures of 10^8 CFU/mL.

2.3. Analysis of MMR gene sequences

Bacterial DNA was extracted and purified using the Qiaamp DNA mini kit (Qiagen, Courtaboeuf, France). The *mutS* (2568 bp), *mutH* (696 bp), *mutL* (1857 bp), *uvrD* (2163 bp), and *dam* (837 bp) genes were amplified by PCR (using primers shown in Table 2) with Phusion high-fidelity DNA polymerase (Finnzymes, Saint Quentin Yvelines, France) according to the manufacturer's recommendations. Amplifications were performed in a PCR express (Thermohybrid) thermocycler with the following parameters; denaturation for 30 s at 98°C, followed by 35 cycles of 10 s at 98°C, 30 s at the specific hybridization temperature, and 45 s at 72°C for polymerization. A final extension step of 10 min at 72°C was included.

Both strands were sequenced in an automatic sequencer (Applied Biosystem 3130 A, Perkin Elmer, Coignières, France) with primers using the Big Dye Terminator Kit version 3.1 (Applied Biosystems, Coignières, France). DNA sequence analysis was carried out using SeqScape software 2.5 (Applied Biosystems, Coignières, France). Nucleotide alignments were determined using the programs ClustalW (<http://www.ebi.ac.uk/clustalw/>) and BLAST (<http://www.ncbi.nlm.nih.gov>, last accessed in November 2007).

2.4. Trans-complementation of the naturally occurring *S. Heidelberg* hypermutator strain

The wild-type *mutS* gene was amplified by PCR using *mutS-SacI* and *SphI* primers (Table 2). The PCR product was digested with the restriction endonucleases *SacI* and *SphI* and ligated into the pGEMT vector (Promega, Charbonnières, France). The vector was transformed into competent cells of *S. Heidelberg* hypermutator (Hm) by heat shock as

previously described (Sambrook et al., 1982). Transformed cells were selectively plated on Tryptic Soy (AES Laboratory, Combourg, France) plates with ampicillin (100µg/mL, Sigma Aldrich, Saint Quentin Fallavier, France).

2.5. 3D representation

S. Typhimurium LT2 and *E. coli* MutS proteins have a high sequence homology (<http://www.ncbi.nlm.nih.gov/blast/Blast.cgi>), which led us to compare their 3D structures. The 3D structure of the *Salmonella* MutS protein was compared to an *in silico* 3D representation of MutS *E.coli* structure (Genbank accession number n°1OH5) (Natrajan et al., 2003), generated using Cn3D software (<http://www.ncbi.nlm.nih.gov/Structure>). This 3D structure was used to analyze the structure of the *Salmonella* MutS protein.

2.6. DNA binding assay

Binding of both wild-type and mutated MutS proteins to DNA was assessed using a gel retardation assay. Proteins from both *S. Heidelberg* wild-type and a hypermutator of the same serovar were expressed in a His₆-tagged vector and extracted and purified using the QIAexpression kit (Qiagen, Courtaboeuf, France), as described below.

To make the His-tagged expression construct, *mutS* DNA from each strain was amplified by PCR using the primers *mutS*-SacI-prot and *mutS*-SphI (Table 2). A dATP was added to the 3' terminus using Taq polymerase enzyme (Promega, Charbonnières, France) in a second and final extension step. Each PCR product was ligated into the pQE-30-UA (ampicillin resistant) vector (QIAexpression kit, Qiagen, Courtaboeuf, France). Recombinant plasmids were transformed into *E. coli* M15 pREP4 (kanamycin resistant) according to the manufacturer's instructions. The orientation of the cloned *mutS* gene was verified by

digestion with *SacI* and *AscI*. In the correct orientation, the *mutS* gene is under the control of PT5 promoter.

His₆-tagged MutS proteins were expressed in *E. coli* cultures incubated overnight in LB broth supplemented with ampicillin (100 µg/mL) and kanamycin (20 µg/mL). Proteins from each strain were extracted (under denaturing conditions), eluted, and purified (under native conditions), according to the QIAexpression kit recommendations. Proteins were purified with a Ni-NTA matrix (Ni-Sepharose 6 Fast-Flow, GE Healthcare, Orsay, France). Protein extracts were mixed with the Ni-NTA matrix for 1 h and then loaded onto a column, proteins were washed with buffer (50 mM NaH₂PO₄, 300 mM NaCl, 20 mM imidazole, pH 7.4) (Sigma Aldrich, Saint Quentin Fallavier, France). Then proteins were eluted in 3 fractions with elution buffer (50 mM NaH₂PO₄, 300 mM NaCl, 250 mM imidazole, pH 7.4) (Sigma Aldrich, Saint Quentin Fallavier, France). Total protein concentration was measured using the 2D quant kit (GE Healthcare, Orsay, France).

DNA from *S. Heidelberg* wild type (wt) and a hypermutator phenotype (Hm) was amplified with primers *RrlA1*, *RrlA2*, and *RrlAbis* (Table 2). The 259 bp PCR products (*rrla1-rrla2* and *rrlabis-rrla2*) were identical, except for a single nucleotide at position 9 (T-C substitution). PCR products were purified using Qiagen columns (Qiagen, Courtaboeuf, France). To prepare heteroduplex DNA, the two DNA fragments were mixed as described by Stanislawska-Sachadyn et al. (2003). The mixture of the two PCR products contained both homoduplex and heteroduplex DNA fragments (Stanislawska-Sachadyn et al., 2003).

The DNA gel retardation assay was performed using 6 and 8 µg of purified His₆-tagged MutS proteins (wt and Hm). The DNA-MutS complexes were prepared as described previously (Stanislawska-Sachadyn et al., 2003). Two controls were used, one with only DNA and one with DNA complexed with BSA (6 µg) (purity 98%, Sigma Aldrich, Saint Quentin Fallavier, France). Each sample was electrophoresed in a 2% TAE agarose gel at 5

mA/cm in a 2% TAE agarose gel, at room temperature for 14 h. Protein-DNA complexes were visualized by staining with ethidium bromide.

2.7. Statistical analysis

Statistical analysis was assessed using 2-sample student's tests or the Wilcoxon 2-sample test using t approximation, where appropriate. Categorical variables were assessed using χ^2 tests exact test. A p value <0.05 was considered to be statistically significant.

3. Results

3.1. Phenotypic identification

No phenotypic differences in identification profiles were observed between *S. Heidelberg* Hm (B182) and wt using the API 20E gallery. Some differences were found using API 32GN; *S. Heidelberg* Hm, in contrast to wt, was not able to grow in the presence of L-alanine, L-serine, lactic acid and propionic acid.

3.2. Mutation rates of *Salmonella* strains

The distribution of rifampicin mutation frequencies of 209 *Salmonella* isolates from veterinary and human origin was determined, using phenotypic categories previously established by Baquero et al. (2004): hypomutable ($f \leq 8 \times 10^{-9}$), normomutable ($8 \times 10^{-9} < f < 4 \times 10^{-8}$), weakly hypermutable ($4 \times 10^{-8} \leq f < 4 \times 10^{-7}$), and strongly hypermutable ($f \geq 4 \times 10^{-7}$) (Figure 1). A significantly higher proportion of hypomutable strains were isolated from humans (38.7%) than from animals (18.4%, $p=0.001$). Among human clinical isolates, only 3.3% were weakly hypermutable compared with strains from animal origin (15.6%, $p=0.011$). Only the B182 bovine strain of *S. Heidelberg* (Hm) (one strain/209, incidence 0.7%) showed a high frequency of mutation using rifampicin, fosfomycin, or nalidixic acid. According to

the criterion described by Leclerc et al. (1996), this strain was categorized as a strong hypermutable isolate. Its frequency of mutation to rifampicin resistance was greater than 50 fold higher (395 ± 100 per 10^8 CFU/mL) than a wild-type strain (wt) of the same serovar (4 ± 2.5 per 10^8 CFU/mL) (Table 3). Its frequency of mutation was also 50 fold higher than the other *Salmonella* strains (median value 2.7 per 10^8 CFU/mL).

3.3. MMR gene sequences of the *S. Heidelberg* hypermutator strain

Sequencing showed no evidence that the MutL, MutH, Dam, and UvrD (CP001120.1) proteins were mutated in the *S. Heidelberg* hypermutable strain B182. Sequences were identical to previous sequenced genes of *Salmonella Heidelberg* SL 476. Mutations observed in the *mutL* gene (Genbank Accession number FJ39011) were silent mutations.

The *mutS* gene of the *S. Heidelberg* B182 (Genbank Accession number FJ539012) was sequenced and analyzed using available databases. A 12 bp deletion in position 1075 to 1086 was identified. In order to determine if the 12 bp deletion might be present in other *Salmonella* serovars and if this deletion constitutes a genetic polymorphism, the sequences of *mutS* genes from several *Salmonella enterica* strains were compared using available databases. Sequence alignments of the *mutS* gene from the genome of *S. Typhimurium* LT2 showed a nearly identical match to all strains of the serovars *Salmonella*: *S. Heidelberg* wt (100%), *S. Choleraesuis*, *S. Paratyphi* A, *S. Typhi* CT18, and *S. Typhi* Ty2 (99%). In contrast, alignment with the *S. Heidelberg* hypermutator, confirmed the 12 bp deletion was only present in this strain (Figure 2).

The nucleotides in this 12 bp sequence encode amino acids LRTA (359-362) (Figure 2), and there were no other modifications in the open reading frame (ORF). The hypothetical location of the LRTA motif on the 3D protein structure of MutS was determined using

<http://www.ncbi.nlm.nih.gov/Structure/CN3D/cn3d.shtml>. The 3D structure was modeled from the *E. coli* MutS protein (E24377A or O157:H7 strain Sakai), which has 95% homology with MutS from *Salmonella* Typhimurium LT2. This analysis revealed the LRTA motif might form a link between 2 beta-sheets in the core domain.

To investigate if the 12 bp deletion could be restored, the *S. Heidelberg* hypermutator strain was trans-complemented with the *mutS* wild-type gene from wt *S. Heidelberg*. The resulting trans-complemented hypermutator strain showed a decrease (~1 log) in the frequency of mutation when exposed to rifampicin, fosfomycin, and nalidixic acid (Table 3). Thus, the wild-type phenotype was partially restored.

3.4. DNA gel retardation assay

The purified proteins MutS Hm and MutS wt were tested in a DNA gel retardation assay for their ability to anchor homoduplex or heteroduplex DNA (with or without a mismatch). The migration of DNA (homoduplexes or heteroduplexes) in the absence of protein on the gel was faster than the migration of DNA in the presence of MutS wt (Figure 3). Therefore, MutS wt protein was able to anchor the DNA (with or without mismatch). Similar results were also obtained for MutS Hm (Figure 3). No binding was observed when DNA was mixed with BSA. Thus, the *mutS*-12 bp deletion does not appear to inhibit MutS from anchoring DNA.

4. Discussion

This work describes the first isolation of a *S. Heidelberg* with a hypermutator phenotype. Hypermutation allows some bacterial cells to adapt to adverse environmental conditions, including antibiotic exposure (Blazquez, 2003; Chopra et al., 2003; Woodford and Ellington,

2007). Stressful conditions, changing environment, or extreme nutrient limitation are known to place bacteria at risk. Bacteria respond through a mutational alteration of genotype (Sundin et al. 2007).

The criteria for distinguishing mutators from non-mutators have been arbitrarily determined by some authors. Besier et al. (2008) demonstrated that thymidine-dependent small-colony variant phenotype was associated with hypermutability in *S. aureus*, but no study was able to show significant phenotypic difference linked to the hypermutator phenotype. Tompkins et al. (2003) determined ~ 0.1% of *E. coli* bacterial cells are hypermutators (with a mutation rate up to 200-fold higher than normal). They also found only 10% of the adaptive Lac + mutations arise in hypermutators. Leclerc et al. (1996) stated cells exhibiting an increase in mutation frequencies of just 50-fold or more should be considered mutators. Accordingly, they reported a higher incidence of mutators, with an average of 3.6% among isolates of *Salmonella enterica* and 1.9% among *E. coli* isolates. Significant variations have been observed in mutation frequencies between clinical isolates from different origins and ecotypes (Baquero et al., 2004; Denamur et al., 2002). Denamur et al. (2002) showed uropathogenic *E. coli* strains had a higher frequency of mutators than enterohemorrhagic and enteroinvasive strains.

Only a few studies are available on the subject of mutation rates of strains isolated from fecal samples. Baquero et al (2004) previously established phenotypic categories, studying a collection of 300 *E. coli* isolates from Spain, and found 69% of fecal strains from healthy volunteers were normo-mutagenic, whereas 11% were weakly mutagenic. They found a higher proportion of hypomutagenic strains (19%) from healthy volunteers than from those with bacteremia or urinary tract infection. The implication of hypermutation in virulence remains conflicting; some authors attesting increasing virulence (Bayliss et al 2008), while others showed hypermutation reduced virulence due to biological cost (Merino

et al. 2002; Montarini et al. 2007). No difference in susceptibility to antimicrobial agents (both quite susceptible, data not shown) was observed between the wild type and hypermutator strain, in agreement Denamur et al (2002) who demonstrated uropathogenic *Escherichia coli* mutator strains were not more resistant than non-mutators. Data on resistance to antimicrobial agents remains controversial in literature (Komp Lingren et al 2003, Baquero et al 2005).

Most mutator strains have mutations in the methyl-directed mismatch repair genes. In the presence of both ATP and a mismatch, MutS recruits MutL. Together they activate MutH and initiate the entire MMR system. The binding of MutS to DNA is very important for successful DNA repair. MutS recognizes errors, such as mismatches, with varying efficiencies (Parker and Marinus, 1992). The crystalline structure of the MutS-DNA complex from *E. coli* has proven very important in understanding the nature of protein-DNA interactions (Mendillo et al., 2007). A mutation in the *mutS* gene causes a hypermutable phenotype (LeClerc et al., 1996; Matic et al., 2006). Leclerc et al. (1996) examined a natural *Escherichia coli* O157:H7 strain and found a 200 bp deletion at the 3' end of the *mutS* gene, which conferred a mutator phenotype. Additionally, they examined five strains of natural populations of *Salmonella enterica* and found three out of five carried a *mutS* mutation. The other two strains had mutations in *mutH* and *uvrD*. In this work, a 12-bp deletion (AA358-362, 1075-1087 bp) was identified in the *mutS* gene of *S. Heidelberg*. A complementation assay indicated the wt *mutS* gene only partially restored repair activity in the *mutS* deficient *S. Heidelberg* Hm. This could indicate a partial activity of MutS Hm or a mutation or dysregulation of regulatory gene(s). DNA mismatch repair is an evolutionarily conserved process that corrects mismatch errors that escape proofreading during DNA replication. Defects in this system usually result in a hypermutable phenotype (Leclerc et al 1996; Matic et al. 2006). DNA repair is initiated upon recognition of a mismatch by MutS. MutS, which

binds to insertion-deletion-loops (IDL's) that are 1- 4 nucleotides in length, also binds to 7 and 8 bp mismatches (Hsieh, 2001; Wang et al., 2003).

The 3D structure of *E. coli* MutS revealed 6 functional domains. From 5' to 3', they are: (1) the mismatch binding site (aa 1 to aa 115), (2) the connector (aa 116 to aa 257), (3) the core domain (aa 266 to aa 443), (4) the lever clamp domain (aa 444 to aa 567), (5) the ATPase domain (aa 568 to aa 765), and (6) the helix-turn-helix (dimer contact) in the C-terminal domain (aa 766 to stop) (Marti et al., 2002). Mutational studies have identified some functions of these domains. Calmann et al. (2005) showed deletion of 53 amino acids from the C-terminus of MutS resulted in (1) an increased mutation rate of about 60-fold and (2) a MutS-protein with a lower affinity than MutS for certain oligonucleotides with base pair mismatches, that in the presence of other MMR components reduced the efficiency of MutH-induced incision at hemimethylated GATC sequences *in vitro*. Miguel et al. (2007) showed the K850A substitution destabilized the tetramer structure which is the active form of MutS. The authors proposed the C-terminal extreme alpha helix (aa 839-850) might play an important role in protein oligomerization. Mendillo et al. (2007) demonstrated dimerization, but not tetramerization, of the MutS C terminus is essential for mismatch repair. Two models have been proposed in *E. coli* to explain the initiation of the mismatch repair system: (1) *cis* model: movement of the MutS or MutS-[MutL and/or other proteins] complex along the helix between the mismatch and the d(GATC)-strand signal and (2) *trans* model: the two sites interact in a DNA-looping mechanism (Kolodner et al., 2007). Assembly of the MutS-MutL heteroduplex complex is sufficient to activate both MutH endonuclease and DNA helicase II. A conformational change in MutS dimers directs recruitment of MutL dimers and might allow the formation of two channels that can receive DNA. The *mutS* 12 bp deletion appeared to have no influence on the ability of the protein to anchor or bind DNA. It is hypothesized the 12 bp deletion in the core domain of MutS may either prevent binding to MutL or modify the

conformation of the MutS dimer. In support of this idea, Joseph et al. (2005) suggested conformational changes in *Haemophilus influenzae* MutS modulate mismatch recognition and trigger some of downstream repair processes. Crystallization of the *Salmonella* MutS protein would reveal if MutS-MutL interactions are present.

Bacterial mismatch repair (MMR) deficient mutants, exhibiting high mutation and homologous recombination rates, are frequently found in natural populations. These high mutation rates are expected to favor their environmental adaptation. The 12 bp deletion in the *mutS* gene of the *S. Heidelberg* Hm was only partially responsible for the hypermutator phenotype observed and has no influence on anchorage of the MutS protein on DNA. Mechanisms other than the MMR system might be involved in hypermutation. A better understanding of the genetic mechanisms and regulatory schemes of hypermutation will help anticipate behavior of hypermutable strains in mixed bacterial populations and their role in survival ability.

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Conflict of interest statement

All authors disclose there are no financial and personal relationships with other people or organizations that could inappropriately influence their work.

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Table 1 Distribution of different serovars of *Salmonella enterica* isolates studied

<i>Salmonella enterica</i> serovars	Sources of the strains					
	Number of isolates					
	Human	Cattle	Sheep	Pig	Poultry	Total
S. Typhimurium	17	58	1	18	7	101
S. Montevideo	1	32	0	0	0	33
S. Enteritidis	15	2	1	0	0	18
S. Hadar	7	0	0	0	0	7
S. Derby	1	2	0	4	0	7
S. Virchow	5	0	0	0	0	5
S. Heidelberg	3	2	0	0	0	5
Other	13	16	1	3	0	33
Total	62	112	3	25	7	209

Table 2 Primers used in this study^a

Primers used for sequencing	Sequences 5'→3'
<i>mutLA</i>	TAT ATA CCG GGG GTT GCG TA
<i>mutLB</i>	TAA ATG AGG GCG GAA TCA AC
<i>mutL1</i>	AGTAAAATCGCCTCGCTTGA
<i>mutL2</i>	GTTTCTCGAACAGGCACTGG
<i>mutL3</i>	GCCCTGCTTCTGATAAC
<i>mutL7</i>	CAAATTCATGCGTACCGAAA
<i>mutL8</i>	CTTTTAAACGCAGCGGAATC
<i>mutSlong1</i>	GAGCATGGTCTGAACTAATTGC
<i>mutSlong2</i>	CCCTTATTTACGCTGTCTGTCT
<i>mutS1</i>	CCCCTGTTGTAGTCCAGGAA
<i>mutS2</i>	GGCTATCTGGCAGGATGGTA
<i>mutS3</i>	GGCCAGGTTCTGGGTAATTT
<i>mutS4</i>	AAGTCGGCTATAACGCGGTA
<i>mutS5</i>	CGGGTAAAAATACGGTCGAT
<i>mutS6</i>	CGACAGTCACAGCCTGAAAA
<i>mutS7</i>	CCAATATGGGCGGTAAAAGT
<i>mutS8</i>	GGTAGCCGAATGCTTAAACG
<i>mutS9</i>	TTTGACCACGGCTAATCTGA
<i>mutS10</i>	GAATTACACCTGCCCCACAT
<i>mutS11</i>	GCTGTTTTATGACGACGACGAA
<i>mutS12</i>	CGCCGTCGCCCGTTGTGGGA
<i>mutS13</i>	TGCTTGACTGTACCGTGACGCC
<i>mutS14</i>	ACGGCGATACTATCGCGTTTAT
<i>mutH1</i>	AGAATTGGAAAACCCGCTTC
<i>mutH4</i>	TGCAAGACTTCGCCATGA
<i>dam1</i>	CTGCAATTGCCTGTGAGTGT
<i>dam4</i>	CGTACTGACGCAGGGATTTC
<i>uvrDA</i>	GTTTACTGCCGCATATCGAA
<i>uvrDB</i>	CGAAACGCGAAAAAGCAATA
<i>uvrD2</i>	AGCGAAGATCAGATGCGTTT
<i>uvrD3</i>	GAGCAGAATTACCGCTCGAC
<i>uvrD4</i>	GCACACCTGCCACAATGTTA
<i>uvrD5</i>	CTGCATGGCACGGGTAAC
Primers used for trans complementation experiment	Sequences 5'→3'
<i>mutS-SacI</i>	GTAAAGAGCTCATGAATGAGTCATTTGATA
<i>mutS-SphI</i>	TCTGGCATGCTTACACCAGACTTTTCAG
Primers used for DNA binding assay	Sequences 5'→3'
<i>rrlA1</i>	TTCTAACGTGGACCCGTAAT
<i>rrlA2</i>	GTTATCCCCGGAGTACCTTT
<i>rrlA bis</i>	TTCTAACGCGGACCCGTAAT
<i>mutS-SacI-prot</i>	AGTAAAGAGCTCATGAATGAGTCATTTGAT

^a Restriction sites are underlined.

Table 3 Mutation rates of the different *S. Heidelberg* used in this study (selection on rifampicin 100µg/ml).

<i>Salmonella Heidelberg</i> strains	Number of mutator strains per 10 ⁸ CFU/mL		
	<i>Rifampicin</i>	<i>Nalidixic acid</i>	<i>Fosfomycin</i>
	100 µg/mL	50 µg/mL	100 µg/mL
Wild-type (wt)	4 ± 2.5	0	28 ± 6.5
B182 hypermutator (Hm)	395 ± 100	236 ± 60	12360 ± 1310
<i>mutS</i> wild-type <i>trans</i> complemented hypermutator	65 ± 20	30 ± 15	1568 ± 650

Figure legends

Figure 1: Distribution of rifampicin mutation frequencies (f) of 209 *Salmonella* isolates from veterinary (dark) and human (white) sources. Categories: hypomutable (hypo) ($f \leq 8 \times 10^{-9}$), normomutable (normo) ($8 \times 10^{-9} < f < 4 \times 10^{-8}$), weakly hypermutable (weak) ($4 \times 10^{-8} \leq f < 4 \times 10^{-7}$), and strongly hypermutable (strong) ($f \geq 4 \times 10^{-7}$).

Figure 2: Partial sequence alignment of the *mutS* gene and MutS protein of four *Salmonella* strains: S.Typhimurium LT2 (GenBank NC0031971), S.Heidelberg wt, SL476 (GenBank CP001120), and S.Heidelberg Hm (B182) (GenBank FJ539011). Deleted nucleic acids and amino acid are shown in bold.

Figure 3: The DNA gel retardation assay. 20 μ L of each sample (His₆-tagged *S. Heidelberg* MutS wt or Hm) was loaded on a 2% gel. Their ability to anchor to homodimer and heterodimer DNA was tested (in lanes 2 to 5). Their ability to anchor to only homodimer DNA was tested (in lanes 7 to 10). The DNA molecular weight marker is in lane 11. Lanes 1 and 6 show the DNA migration in the absence of added MutS protein. MW: Molecular weight marker VIII (Roche, Meylan, France).

Figure 1

Figure 2

<i>S.</i> Typhimurium LT2	GCG	<u>CTG CGC ACC GCG CGT</u>	CCG
	A	L R T A R	P
<i>S.</i> Heidelberg wt	GCG	<u>CTG CGC ACC GCG CGT</u>	CCG
	A	L R T A R	P
<i>S.</i> Heidelberg SL476	GCG	<u>CTG CGC ACC GCG CGT</u>	CCG
	A	L R T A R	P
<i>S.</i> Heidelberg Hm (B182)	GCG	C-----GT	CCG
	A	R	P

Figure 3.