**S1 Table. Strains and plasmids used in this study**

|  |  |  |
| --- | --- | --- |
| Strain or plasmid | Relevant characteristics | Reference |
| Strains |  |  |
|  *E. coli* |  |  |
|  DH5α™ | F– Φ80Δl*acZ*Δ*M15* Δ(*lacZYA*-*argF*) U169 *recA1* *endA1* *hsdR17* (rK–, mK+) *phoA* supE44 λ– *thi-1* *gyrA96* *relA1* | Invitrogen™ |
|  *S. agalactiae* |  |  |
| BM110 | Serotype III, ST-17, human clinical isolate | [30] |
| BM110*covR* | In frame deletion of *covR* in BM110 | Asma Tazi |
| BM110CovRD53A | BM110 mutant expressing CovR with a D53A substitution | Arnaud Firon |
|  BM110*∆bp* | In frame deletion of *sbp1* (*san1519 according to COH1*) | This study |
|  BM110*∆bp* bWT | Back to the WT strain obtained during the construction of BM110*∆bp* | This study |
|  BM110*∆43* | Deletion of the 43-bp sequence (5' GTTTTAAATAATAAAAAAAGCCATATATCAATTTGATATATGGC) | This study |
|  BM110*∆43* bWT | Back to the WT strain obtained during the construction of BM110*∆43* | This study |
| A909 | Serotype Ia, ST-7, human clinical isolate | [31] |
|  A909*∆bp* | In frame deletion of *bp* (*sak1439*) | This study |
|  A909*∆bp* bWT | Back to the WT strain obtained during the construction of A909*∆bp* | This study |
| NEM316 | Serotype III, ST-23, human clinical isolate | [32] |
| NEM316*covR* | In frame deletion of *covR* in NEM316 | [33] |
| NEM316CovRD53A | NEM316 mutant expressing CovR with a D53A substitution | [33] |
|  *L. lactis* |  |  |
|  NZ9000 | *L. lactis* subsp. *cremoris* MG1363 containing the *nisRK* gene in the genome | [34] |
| Plasmids |  |  |
|  pTCV*gfp* | *EGFP* expression vector  |  |
|  pTCV1 | pTCV*gfp* with 125-bp PCR product ( 2bUp14a / 2bUp12) from A909b | This study |
|  pTCV2 | pTCV*gfp* with 356-bp PCR product ( 2bUp6 / 2bUp20) from A909 | This study |
|  pTCV3 | pTCV*gfp* with 406-bp PCR product with 2bUp6 / 2bUp12 from A909 | This study |
|  pTCV4 | pTCV*gfp* with 38-bp DNA fragment (2bUp1/2bUp2) (AAACGATAATTTAAGGTTCAGTTAAGGAAGTAATCGCG) | This study |

a Primer sequences are listed in Table S2.

b Genomic DNA of the corresponding strain was used as template for sequence amplification