

Table S1. Bacterial strains and plasmids

Strains and plasmids	Description	Reference
<i>Escherichia coli</i>		
DH5 α	F- ϕ 80lacZ Δ (lacZYA-argF) U169 deoRsupE44 Δ lacU169 (f80lacZDM15) hsdR17 recA1 endA1 (rk- mk+) supE44gyrA96 thi-1 gyrA69 relA1	(7)
BL21(DE3) <i>plysS</i>	F- ompT hsdS gal (rb- mb+) DE3(Sam7 Δ nin5 lacUV5- T7 Gen1)	(7)
BL21(DE3) <i>plysS</i> pET11b- <i>gbaA</i>	For overexpression of His-tagged GbaA	This study
BL21(DE3) <i>plysS</i> pET11b- <i>gbaAC55S</i>	For overexpression of His-tagged GbaAC55S	This study
BL21(DE3) <i>plysS</i> pET11b- <i>gbaAC104S</i>	For overexpression of His-tagged GbaAC104S	This study
<i>Staphylococcus aureus</i>		
RN4220	restriction negative strain/MSSA cloning intermediate derived from 8325-4	(3)
COL	Archaic HA-MRSA strain	(6)
COL- Δ <i>gbaA</i>	COL <i>gbaA</i> deletion mutant	This study
COL- Δ <i>gbaB</i>	COL <i>gbaB</i> deletion mutant	This study
COL- Δ <i>gbaA</i> -pRB473- <i>gbaA</i>		This study
COL- Δ <i>gbaA</i> -pRB473- <i>gbaAC55S</i>		This study
COL- Δ <i>gbaA</i> -pRB473- <i>gbaAC104S</i>		This study
COL- Δ <i>gbaB</i> -pRB473- <i>gbaB</i>		This study
COL- Δ SACOL2590-92	COL SACOL2590-92 deletion mutant	This study
<i>Staphylococcus</i> phage 81		
pET11b	<i>E. coli</i> expression plasmid	Novagen
pRB473	pRB373-derivative, <i>E. coli</i> / <i>S. aureus</i> shuttle vector, Amp ^r , Cm ^r	(1)
pRB473-XylR	pRB373-derivative, <i>E. coli</i> / <i>S. aureus</i> shuttle vector, containing xylose-inducible P _{Xyl} promoter Amp ^r , Cm ^r	(4)
pET11b- <i>gbaA</i>	pET11b-derivative for overexpression of His-tagged GbaA	This study
pET11b- <i>gbaAC55S</i>	pET11b-derivative for overexpression of His-tagged GbaAC55S	This study
pET11b- <i>gbaAC104S</i>	pET11b-derivative for overexpression of His-tagged GbaAC104S	This study
pMAD- <i>gbaA</i> deletion	pMAD with up- and downstream region of <i>gbaA</i>	This study
pMAD- <i>gbaB</i> deletion	pMAD up- and downstream region of <i>gbaB</i>	This study
pMAD-SACOL2590-92 deletion	pMAD with upstream region of SACOL2592 and downstream region of SACOL2590	This study
pRB473-XylR- <i>gbaA</i>	pRB473-derivative expressing <i>gbaA-His</i> under P _{Xyl}	This study
pRB473-XylR- <i>gbaAC55S</i>	pRB473-derivative expressing <i>gbaAC55S-His</i> under P _{Xyl}	This study
pRB473-XylR- <i>gbaAC104S</i>	pRB473-derivative expressing <i>gbaAC104S-His</i> under P _{Xyl}	This study
pRB473-XylR- <i>gbaB</i>	pRB473-derivative expressing <i>gbaB</i> under P _{Xyl}	This study

Table S2. Oligonucleotide primers

Primer name	Sequence (5' to 3')
pET-gbaA-for-NheI	CTAGCTAGCATGCGAAAAGATGCAAAAGAGA
pET-gbaA-rev-BamHI	CGCGGATCCTTAGTGATGGTGATGGTGATG-GTCATTACGTCCCACCTCAT
gbaAC55S-for	GATAAAAGCGATTTAT CT TACTACGTCATACAA
gbaAC55S-rev	TTGTATGACGTAGTA AG ATAAATCGCTTTTATC
gbaAC104S-for	AAGGCACTACTGCAAT CT TATTGAAGCAGGCAAC
gbaAC104S-rev	GTTGCCTGCTTCAAT AG ATTGCAGTAGTGCCTT
pMAD-gbaA-for-BglII	CGCAGATCTAACTGCATTACCTTGCTTCC
pMAD-gbaA-f1-rev	TAATTTCTGTACGCTTCAATTGTGCATCTTTTCGCATGATTACA
pMAD-gbaA-f2-for	TGTAATCATGCGAAAAGATGCACAATTGAAGCGTACAGAAATTA
pMAD-gbaA-rev-Sall:	CCAGTCGACGCATTGTTACTGCCGATTTAG
pRB-gbaA-for-BamHI	TAGGGATCCGAATTTTAAGTAGGTGTAATCATGCGAAAAGATGC AAAAGAGA
pRB-gbaA-His-rev-KpnI	CTCGGTACCTTAGTGATGGTGATGGTGATGGTCATTACGTCCCA CCTCAT
pMAD-gbaB-for-BglII	CGCAGATCTCAAACCTTTTGGATGAAGAAGGC
pMAD-gbaB-f1-rev	TCTCCATCGCCATTAATAAATGTCTGCACTTGCATAGCCTAA
pMAD-gbaB-f2-for	TTAGGCTATGCAAGTGCAGACATTTAATAATGGCGATGGAGA
pMAD-gbaB-rev-Sall	CCAGTCGACATCCTAAATCATTGATGCAACG
pRB-gbaB-for-BamHI	TAGGGATCCATTTAGATGAGGTGGGACGTA
pRB-gbaB-rev-KpnI	CTCGGTACCCTACCAAGGCATCTCTCCAT
pMAD-SACOL2590-92-for-BglII	CGCAGATCTTTCTTGATAAAATGCGCTTTGG
pMAD- SACOL2590-92-f1-rev	GAGAATTTCAATTTAGCAGACCATAAGGTAGATGCTTCGAAATG
pMAD- SACOL2590-92-f2-for	CATTTCGAAGCATCTACCTTATGGTCTGCTAAAATGAAATTCTC
pMAD- SACOL2590-92-rev-Sall	CCAGTCGACATCAACACTTGATTCTAAGGAG
Emsa-gbaA-for	TATCAACACTCTTTTCTTTTATG
Emsa-gbaA-rev	TCTTTTGCATCTTTTTCGCATGA
NB-gbaB-for	TCACAGGAGGCAATAAAGGGT
NB-gbaB-rev	CTAATACGACTCACTATAGGGAGAATTTGTAGCGCCTGGATCAG
NB-SACOL2590-for	GCAAGATTTAGAAAAGAGCGCA
NB-SACOL2590-rev	CTAATACGACTCACTATAGGGAGAACCCTTGGCTTACTGTTGGT

Restriction sites are underlined and bold bases indicate point mutations.

Supplementary References

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