

Supplementary Material

Table S1. Primers for PCR analysis

Number	Primer sequence	Position
Pyruvate dehydrogenase E2 component(PDHE2)		
E2-F	<u>GGATCC</u> CATGTTAAAATGCAATTACAGACG	1→25
E2-R	CTCGAGTTAGTTGTTCAATGTGAAAAATT	212←237
Purine-nucleoside phosphorylase(PNP)		
PNP-F	<u>GGATCC</u> CATGACACCACACATAAATGCACC	1→23
PNP-R	CTCGAGTTATGCTAACTCTAAAGCTATTT	677←702
glycerol ABC transporter(UgpB)		
ugpB-1F	<u>GGATCC</u> CATGTTGAAAAAAATTATTAAAGCTG	1→28
ugpB-1R	GCTTTAAAGCATAACCAAAATTGATTGTT	679←707
ugpB-2F	GAACAATCAATTGGTATGCTTAAAGC	679→707
ugpB-2R	GAATTATTGTTAGTCTTCATAATTCTTGG	999←1028
ugpB-3F	CCAAGAATTAT <u>GG</u> AAAGACTACAATAATT	999→1028
ugpB-3R	AGATTGCCATGAACCCCATTCTTTG	1115←1141
ugpB-4F	CAAAAGAAC <u>GGGG</u> TGATGGCAAATCT	1115→1141
ugpB-4R	TGTACCAGTTATCTTCTTCGCCAGAACCTTGTT TTGCCTGTGTATATCCATT	1416←1474
ugpB-5F	TAAATGGATATA <u>ACACAGG</u> AAAAACAAGGTTCTG GCGAAGAAGAAGATAACT <u>GG</u> TACA	1416→1474
ugpB-5R	<u>CTCGAG</u> TTATTGTTTTAGTTGTTCAAATG	1853←1857
membrane lipoprotein P81		
P81-1F	<u>GGATCC</u> ATGAGTAAGAAAAATAAATTAAATGATTG	1→28
P81-1R	AATGAAGCCATAGTATTCCATTGTGGCT	155←182
P81-2F	AGCCACAAT <u>GG</u> AATACTATGGCTTCATT	155→182
P81-2R	TTTCAAATT <u>CACCC</u> CATAATTGGTACT	675←703
P81-3F	AGTACAAAAATTAT <u>GGGG</u> GAATTGAAA	675→703
P81-3R	TCTTAAGAATA <u>AGCC</u> ATTGTTATTATC	1441←1470
P81-4F	GATAATAAAC <u>ATGG</u> CATTATTCTTAAGA	1441→1470
P81-4R	TCTTTCATTAGATTATT <u>CC</u> ATTACCAAG	1664←1693
P81-5F	CTGGTAA <u>ATGG</u> AATAATCTAATGAAAAGA	1664→1693
P81-5R	<u>CTCGAG</u> TTATTAAAGAATTGACTGCTTGATG	2162←2187

Underline indicates introduced restriction enzyme sites. Bold indicates TGA-to-TGG changes of the codon for permitting tryptophan expression in *E. coli*.

Table S2. Association between color change and mycoplasma growth time.

Fresh normal RS severed as a source of complement, and samples of *M. bovis* were grown in complete PPLO broth medium containing one of various rabbit antisrum. The color of the medium, which changes with *M. bovis* growth, was divided into categories corresponding to four growth stages.

NC	PC	NS		anti- <i>M. bovis</i>		anti-PDHE2		anti-PNP		anti-P81		anti-UgpB	
		RS	HI	RS	HI	RS	HI	RS	HI	RS	HI	RS	HI
0h	-	-	-	-	-	-	-	-	-	-	-	-	-
12h	-	+	-	+	-	+	-	+	-	+	-	+	-
24h	-	++	+	++	-	++	+	++	+	++	-	++	-
36h	-	+++	++	+++	-	+++	++	+++	++	+++	-	+++	-
48h	-	+++	+++	+++	-	+++	+++	+++	+++	+++	-	+++	-

NC:negative control,2mL PPLO broth; PC: positive control, 2mL PPLO broth containing 5×10^4 CFU/mL *M. bovis*. RS: with normal rabbit serum as the complement source; HI: with heat-inactivated normal rabbit serum as control. Grade of color: -, red; +,yellowish red; ++, orange; +++, yellow.

Figure Captions

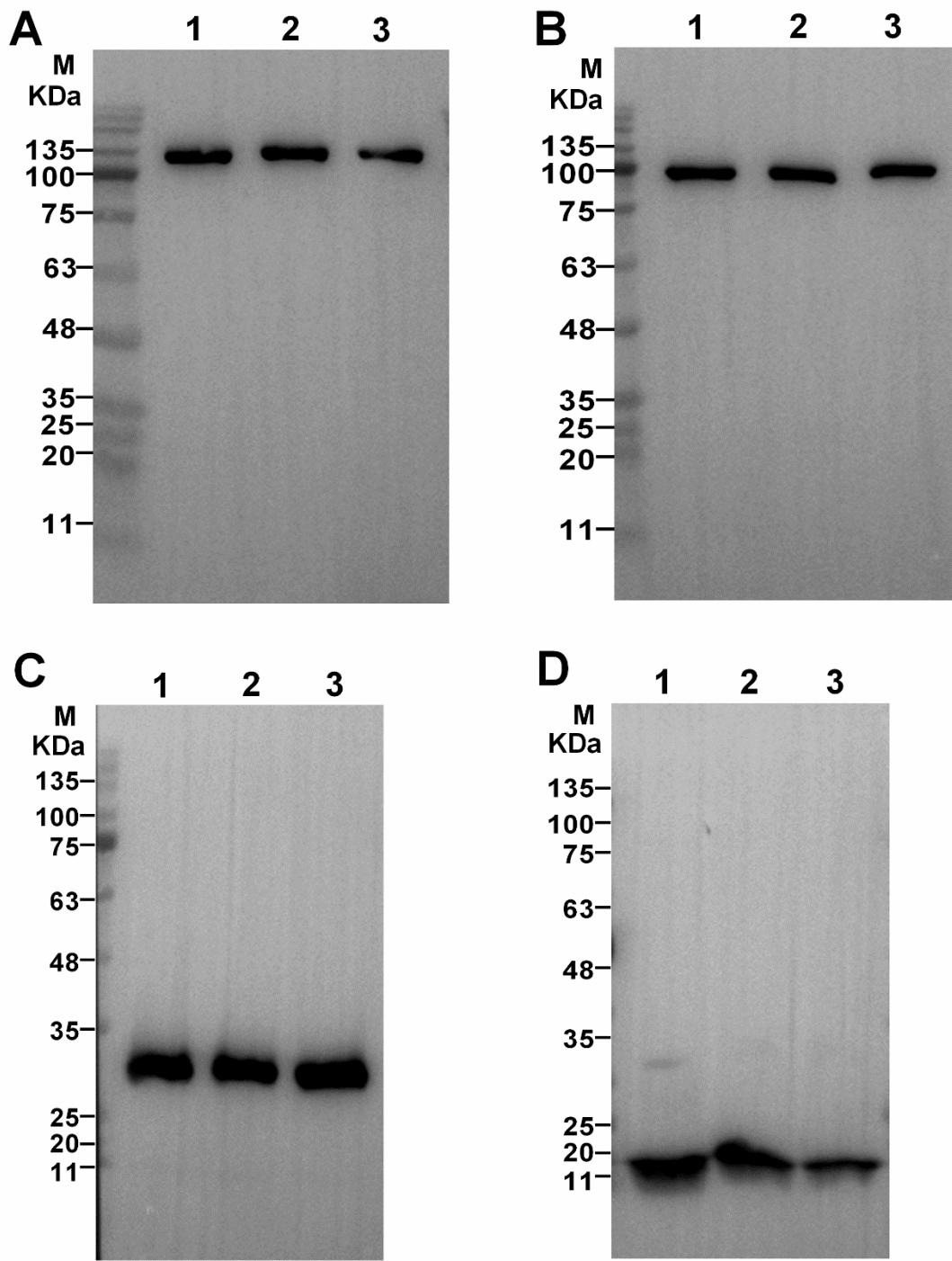


Fig.S1. Western blotting analysis. Western blotting was conducted with 25 µg of whole cell proteins from *M. bovis* PG45 (Lanes 1), PD (Lanes 2), HB (Lanes 3), using rabbit anti-P81 pAb (A), rabbit anti-UgpB pAb (B), rabbit anti-PNP pAb (C), rabbit anti-PDHE2 pAb (D) as the primary antibody, respectively.