Table S1. Primers and conditions of amplification used for *Gordonia* species.

Gene	Sequence *(5´-3´)*	Size (bp)	Reference
16S rDNA amplification		1,459	[57]
fD1: forward primer	AGAGTTTGATCCTGGCTCAG		
rP2: reverse primer	ACGGCTACCTTGTTACGACTT		
16S rDNA sequencing			[58]
fD1 and rP2			
E781: forward primer	GATTAGATACCCTGGTAG		
U1115: reverse primer	GGGTTGCGCTCGTTG		
*sec*A1 amplification and sequencing		496	[23]
SecA1-F: forward primer	GTAAAACGACGGCCAGGACAGYGAGTGGATGGGYCGSGTGCACCG		
SecA1-R: reverse primer	CAGGAAACAGCTATGACGCGGACGATGTAGTCCTTGTC´		
*gyr*B amplification and sequencing		1,230	[23]
UP-1: forward primer	GAGGTCGTCATGACCCAGCTGCAYGCNGGNGGNAARTTYGA		
UP-2: reverse primer	*AGCAGCGTCGAGATGTGCTGGCCRTCNACRGCNGCRTCNGTCA*		
*Gordonia bronchialis gyr*B amplification and sequencing		1,125	This study
123F: forward primer	GAGGTCGTCATGACCCAGCTGCAYGCNGGNGGNAARTTYGA		
1248Rev: reverse primer	*AGCAGCGTCGAGATGTGCTGGCCRTCNACRGCNGCRTCNGTCA*		

The amplification reaction was performed under the following conditions: initial denaturation at 94^{0}C for 5 min followed by 35 cycles of denaturation at 94^{0}C for 30 s, primer annealing at 55^{0}C for 30 s, and primer extension at 72^{0}C for 1 min with final extension at 72^{0}C for 10 min. The temperature of annealing of 55^{0}C, was diminished to obtain amplification band (50^{0}C, 46^{0}C).