

Supplementary materials

<https://doi.org/10.1631/jzus.B2000606>

Development of a *pyrF*-based counterselectable system for targeted gene deletion in *Streptomyces rimosus*

Yiying YANG^{1*}, Qingqing SUN^{2*}, Yang LIU¹, Hanzhi YIN², Wenping YANG¹, Yang WANG¹, Ying LIU¹, Yuxian LI¹, Shen PANG², Wenxi LIU², Qian ZHANG², Fang YUAN², Shiwen QIU², Jiong LI³, Xuefeng WANG³, Keqiang FAN², Weishan WANG², Zilong LI²✉, Shouliang YIN¹✉

Table S1 Similarity and identity values of orotidine-5'-phosphate decarboxylase amino acid sequences among *Streptomyces* strains

| Source | Amino acid residues | GenBank accession No. | Similarity | Identity |
|------------------------|---------------------|-----------------------|------------|----------|
| <i>S. rimosus</i> | 286 | ELQ81494.1 | -- | -- |
| <i>S. avermitilis</i> | 280 | KUN53591.1 | 81.2% | 77.7% |
| <i>S. coelicolor</i> | 281 | TYP05411.1 | 86.1% | 90.4% |
| <i>S. venezuelae</i> | 278 | WP_150188059.1 | 81.8% | 76.9% |
| <i>S. aureofaciens</i> | 277 | WP_030290267.1 | 82.1% | 74.3% |

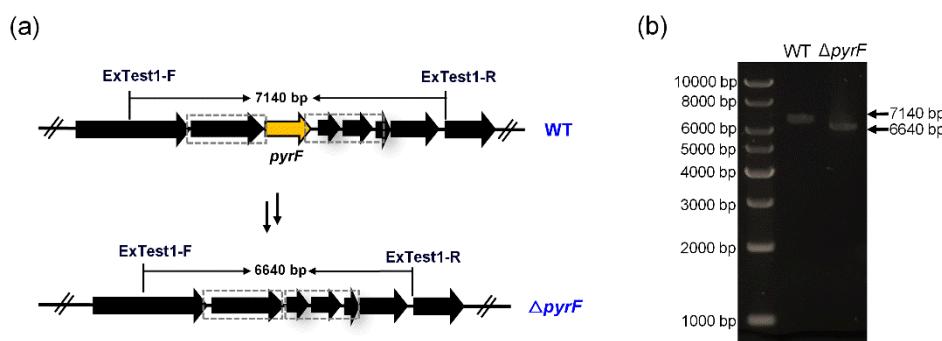


Fig. S1 The Δ pyrF strain was identified by PCR amplification using external primers ExTest1-F/ExTest1-R (primers found only in the genome), the expected amplicon for WT strain is 7140 bp and that 6640 bp for the Δ pyrF strain.

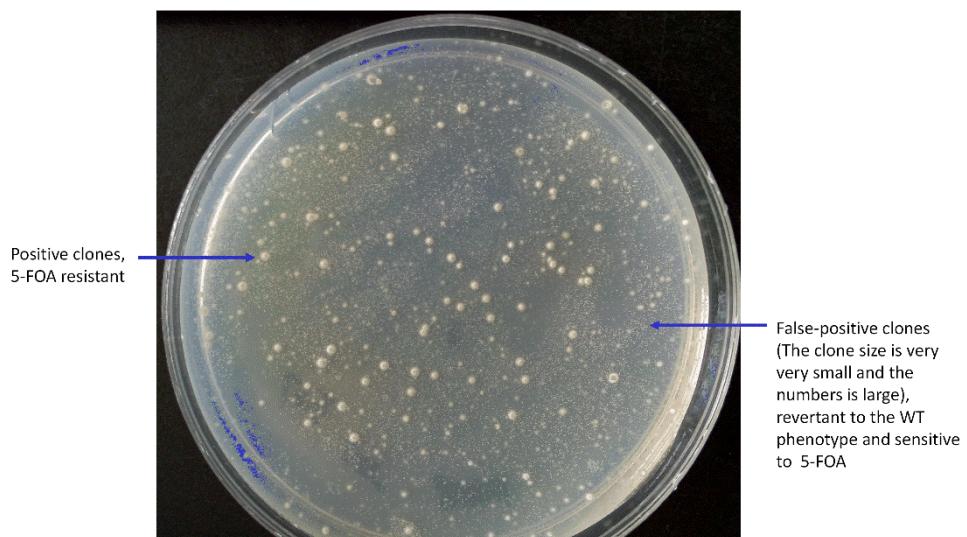


Fig. S2 Characteristics and phenotype of the $\Delta pyrF \Delta tcr$ strain screened on solid MM medium with 5-FOA (100 μ M) and uracil (300 μ M).