

Fig. S1 Scanning electron micrograph of *T. pulmonis* TP-B0596 (a), the liquid culture of *S. lividans* with *T. pulmonis* liquid-culture broth sterilized in an autoclave (b), Co-culture with regenerate cellulose membrane in a dialysis flask (c, d), and the addition of *T. pulmonis* cell wall fraction into *S. lividans* pure culture (e).

(a) Scanning electron micrograph of *T. pulmonis* TP-B0596. The bar represents 1 μm. (b) The liquid culture of *S. lividans* with *T. pulmonis* liquid-culture broth sterilized in an autoclave at 121°C for 0, 1, 3, 10, or 30 min. The cultures were grown at 30 °C for 7 days. (c) Co-culture of *T. pulmonis* and *S. lividans* in a dialysis flask without dialysis membrane (positive control). (d) Co-culture of *T. pulmonis* and *S. lividans* in a dialysis flask with regenerated cellulose membrane (pore size is about 50 kDa). *T. pulmonis* was grown in the left compartment and *S. lividans* in the right compartment at 30 °C for 7 days. (e) The addition of cell wall fraction into *S. lividans* pure culture (right), and no addition (left) at 30 °C for 7 days. The cell wall was prepared from a 500-ml culture of *T. pulmonis* pure culture.



Fig. S2 Co-culture of widely divergent microorganisms and *S. lividans*

(a) The combined cultures of *S. lividans* and *Tsukamurella* strains were grown at 30 °C for 7 days. The left side of the K-1 flask contains the pure culture of *S. lividans*. Red-pigment production was induced in the combined cultures of *S. lividans* with *T. pseudospumae*, *T. spumae* and *T. strandjordii*. (b) Widely divergent microorganisms and *S. lividans* were co-cultivated in A-3M medium at 30 °C for 7 days. The left side of the K-1 flask contains the pure culture of *T. pulmonis*.

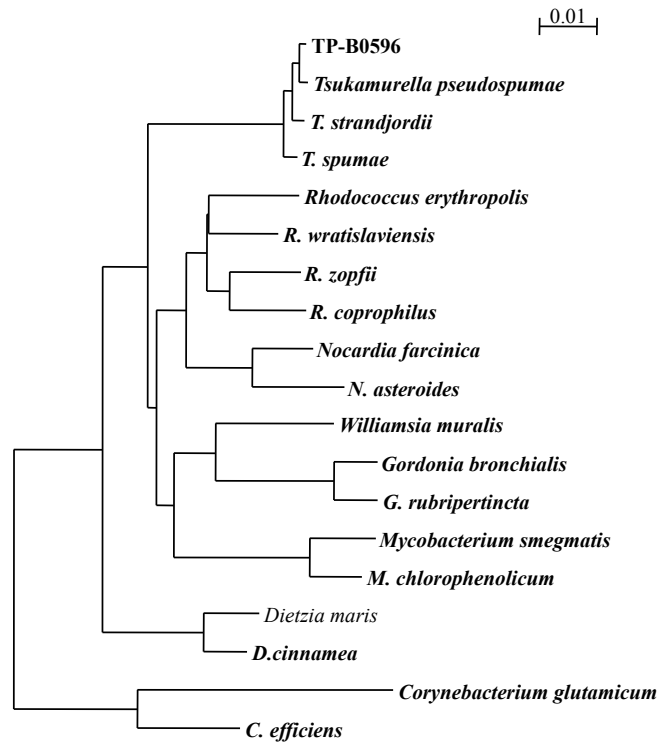


Fig. S3 Phylogenetic tree based on 16S rRNA sequences of microorganisms used in this study.
 Species names in the bold type induced red-pigment formation in *S. lividans*. The bar represents 1% sequence divergence.

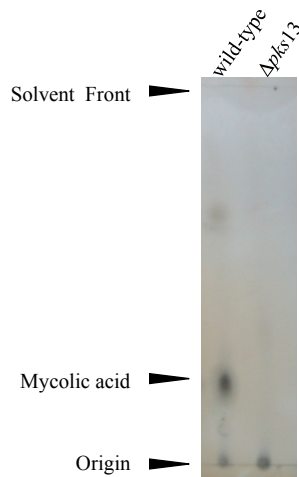


Fig. S4 Mycolic acid detection in *C. glutamicum* wild-type and $\Delta pks13$ strains
 Mycolic acid was extracted from *C. glutamicum* wild-type and $\Delta pks13$ strains that were cultured in LB broth for 2 days. Methyl-esters of the extracted mycolic acids were subjected to thin-layer chromatography (TLC), and the plates were developed in *n*-hexane-diethylether (4:1) and soaked in 50% H₂SO₄.

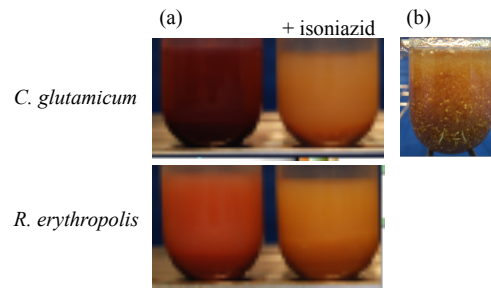


Fig. S5 Addition of mycolic acid synthase inhibitors to co-culture of *S. lividans* with activator strains

(a) Addition of mycolic acid biosynthetic inhibitor isoniazid to the co-culture of *S. lividans* with activator strains. Co-culture of *S. lividans* with *C. glutamicum* and *R. erythropolis* did not induce red-pigment production. (b) Addition of extracted mycolic acid of *C. glutamicum* to a pure culture of *S. lividans* did not induce red-pigment production.

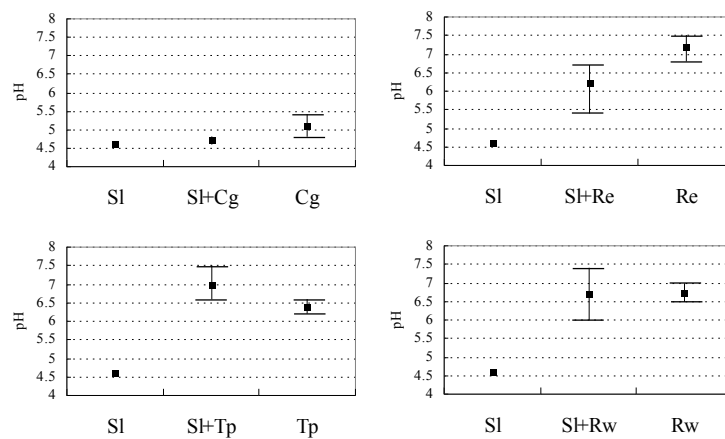


Fig. S6 Measurement of pH of the whole broth of pure cultures and co-cultures.

The pH values of both pure and co-cultures were measured 3 times and the average values were plotted. Cg, Re, Tp, Rw, and SI are abbreviations for *C. glutamicum*, *R. erythropolis*, *T. pulmonis*, *R. wrastrivaensis*, and *S. lividans* and indicate pure cultures of these microorganisms. Cg + SI, Re + SI, Tp + SI, and Rw + SI indicate co-cultures. The cultures were grown in A-3M medium at 30 °C for 7 days.

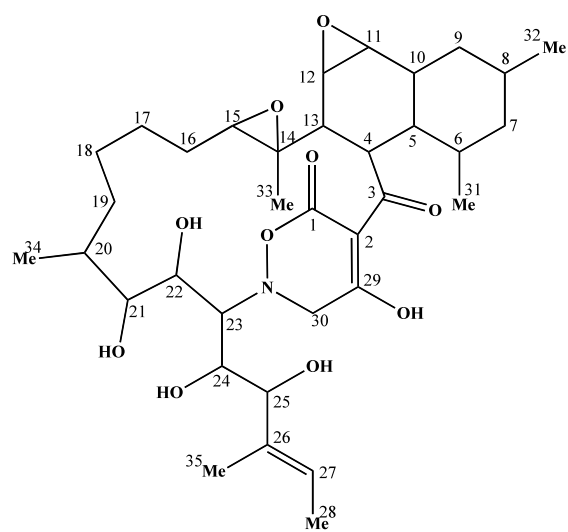


Fig. S7 Determination of carbon positions in alchivemycin A
The numbers represent the carbon positions in alchivemycin A.

Table S1
Nuclear magnetic resonance assignment data of Alchivemycin A

| Position | C (ppm) | H (ppm) | Type |
|----------|---------|--------------------------------|-----------------|
| 1 | 176.61 | | C |
| 2 | 105.28 | | C |
| 3 | 201.30 | | C |
| 4 | 36.79 | 4.72 (t, 10.9) | CH |
| 5 | 38.72 | 2.25 (dm, 11.5) | CH |
| 6 | 39.35 | 1.67 (m) | CH |
| 7 | 39.89 | 1.23 (m), 1.34 (q, 12.7) | CH ₂ |
| 8 | 34.68 | 1.58 (m) | CH |
| 9 | 34.00 | 1.17 (q, 12.4), 1.67 (m) | CH ₂ |
| 10 | 40.36 | 2.19 (dm, 12.9) | CH |
| 11 | 58.70 | 3.11 (m) | CH |
| 12 | 54.54 | 3.05 (d, 3.0) | CH |
| 13 | 51.29 | 2.09 (d, 10.1) | CH |
| 14 | 61.60 | | C |
| 15 | 64.78 | 2.74 (d, 9.6) | CH |
| 16 | 27.44 | 1.47 (m), 1.71 (m) | CH ₂ |
| 17 | 26.54 | 1.72 (m) | CH ₂ |
| 18 | 28.03 | 1.33 (m), 1.58 (m) | CH ₂ |
| 19 | 34.72 | 1.57 (m), 1.62 (m) | CH ₂ |
| 20 | 36.16 | 1.67 (m) | CH |
| 21 | 79.45 | 3.50 (dd, 1.4, 8.8) | CH |
| 22 | 75.32 | 4.30 (dd, 3.8, 8.7) | CH |
| 23 | 62.74 | 3.77 (dd, 1.9, 3.8) | CH |
| 24 | 71.76 | 4.42 (dd, 1.7, 8.1) | CH |
| 25 | 80.38 | 4.06 (d, 8.0) | CH |
| 26 | 137.37 | | C |
| 27 | 124.11 | 5.61 (q, 6.7) | CH |
| 28 | 13.77 | 1.66 (d, 6.8) | CH ₃ |
| 29 | 192.23 | | C |
| 30 | 64.08 | 4.16 (d, 17.0), 4.31 (d, 17.1) | CH ₂ |
| 31 | 22.07 | 0.81 (d, 7.3) | CH ₃ |
| 32 | 23.25 | 1.02 (d, 6.5) | CH ₃ |
| 33 | 15.30 | 1.28 (s) | CH ₃ |
| 34 | 12.67 | 0.83 (d, 7.0) | CH ₃ |
| 35 | 12.14 | 1.68 (s) | CH ₃ |

Table S2, The sample sources of isolated strains

| Bacterial strain | Source of isolation | reference |
|--|---|-----------|
| <i>C. glutamicum</i> ATCC 13869 | nature | (10) |
| <i>C. efficiens</i> NBRC 100395 ^T | soil | (2) |
| <i>R. erythropolis</i> JCM3201 ^T | soil | (4) |
| <i>R. coprophilus</i> JCM3200 ^T | lake mud | (1) |
| <i>R. wratislaviensis</i> JCM9689 ^T | soil | (5) |
| <i>R. zopfii</i> JCM9919 ^T | toluene-phenol bioreactor | (14) |
| <i>T. pulmonis</i> TP-B0596 | soil | This work |
| <i>T. pseudospumae</i> JCM13375 ^T | activated sludge foam | (12) |
| <i>T. spumae</i> JCM12608 ^T | activated sewage sludge foam | (11) |
| <i>T. strandjordii</i> JCM11487 ^T | blood from a 5-year-old girl with acute myelogenous leukemia | (9) |
| <i>G. rubripertincta</i> JCM3204 ^T | ND | |
| <i>G. bronchialis</i> NBRC 16047 ^T | sputa of patients with pulmonary disease | (4) |
| <i>N. farcinica</i> JCM3088 ^T | ND | |
| <i>N. asteroides</i> NBRC 15531 ^T | soil or human infection | (3) |
| <i>M. smegmatis</i> NBRC 3082 | syphilitic chancres | (7) |
| <i>M. chlorophenicum</i> NBRC 15527 ^T | chlorophenol-contaminated lake sediment | (6) |
| <i>W. muralis</i> JCM10676 ^T | Non-water-damaged building material of a children's day care center | (8) |
| <i>D. cinnamea</i> JCM13663 ^T | perianal swab of a human with a bone marrow transplant | (15) |
| <i>D. maris</i> JCM6166 ^T | soil | (13) |

ND : Not described

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