

Modified by Gunther Stier from a protocol described in H. Inoue, H. Nojima, and H. Okayama. "High efficiency transformation of Escherichia coli with plasmids" Gene 96 (1):23-28, 1990.

Procedure

1. Thaw DMSO competent cells on ice. Try to use them as quick as possible, once they thawed!
2. Add 100ng of circular plasmid to 50 μ l cells
3. Incubate 10-30 minutes on ice
4. Heat shock 45seconds at 42°C
5. Chill on ice for 2min
6. Spin down (15sec at 5,000xg) and remove SN [This seems to boost efficiency by a factor of 10, by removing DMSO and Mn salts]
7. Resuspend in 0.2-1.0ml LB or SOC and grow in the shaker 20-60minutes
8. Plate 20 μ l and the rest on plates with the appropriate antibiotic

Needed

1. Heating bath at 42°C
2. Plates prewarmed at 37°C
3. LB or SOC media prewarmed at 37°C