Quick-Screening

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1. Take a single colony of bacteria, draw a 1cm*1cm square on the plate and culture overnight.

2. Take about half of the bacteria:
   + 100μl Lysis buffer (200mM Tris, 3%SDS, pH~12.5)
   Stir the buffer with wood applicator and wait for 15min
   + 100μl phenol/chloroform and invert 50 times

3. Centrifuge at 12000g, 5 min at room temp.

4. Take 5μl supernatant to run gel (don't forget to run the plasmid vector at the same time!).