High Yield Preparation of Genomic DNA from Streptomyces Summary Modified 9.5.2016 by Keith Yamada

- 1. Culture in 30ml **GYM buffer** + **0.5% glycine** + 1.5ml stock culture
- 2. Incubate for 46-96 hrs with shaking at 30°C
- 3. Centrifuge for 5 min at 4000g (optional: store pellet at -20°C)
- 4. Wash 2 x 10ml of **10% sucrose**
- 5. Resuspend in 10ml of lysis solution (+RNase) in a 50ml Falcon tube
- 6. Add 10mg 20mg **lysozyme** + 5mg achromopeptidase
- 7. Incubate for 20 min 40 min at 37°C
- 8. Add 1ml **10% SDS** + 5mg proteinase K
- 9. Incubate for 1.5 hrs at 55°C
- 10. Add 3.6ml **5M NaCl** + 15ml **chloroform**
- 11. Rotate end-over-end for 20 min at 6 rpm
- 12. Centrifuge for 20 min at 5000g 4000g
- 13. Transfer aqueous phase with wide pipet into a clean tube
- 14. Add 1 vol **isopropanol** to precipitate DNA
- 15. Spool using a sealed Pasteur pipet and transfer to a microcentrifuge tube
- 16. Rinse with 1ml cold 70% ethanol
- 17. Air dry DNA
- 18. Dissolve with minimal prewarmed (60°C) 10mM Tris-HCl buffer or MQ H20

Lysis Solution

0.3M sucrose 20.54g C1V1 = C2V2 == V1 = C2V2/C1

25mM EDTA 10ml (500mM) 25mM Tris-HCl 10ml (500mM)

pH 7.5 Fill to 200ml with H20 2 U RNase (add immediately before use)

10% Sucrose Solution

20g / 200ml