

JUNE

Date: Wednesday, 6/27

Objective(s):

Plasmid extraction and PCR to amplify DNA fragments in preparation for plasmid construction.

Procedure:

1. Overnight cultures of DH5 α bacteria carrying the relevant plasmids were spun down at 6400 rpm for 6 min. The supernatant was poured away.
2. The TOM15b MCS and PAL - 4CL PEaaa+5 backbones were purified from the cell pellet. Refer to the protocol '[Plasmid DNA Purification Using QIAprep Spin Miniprep Kit and a Microcentrifuge](#)'.
3. The concentration of the purified plasmids was measured using NanoDrop.
4. PCR was carried out to amplify enzymes

Sample	Concentration (ng/ μ L)
MCS - OsPKS TOM15b	120
PAL - 4CL PEaaa+5	32