Exonuclease I/ Phosphatase

Exo I/Phos 10 x buffer

Stock concentrations	volume (ml)	Final concentration
1M MgCl2	5 ml	100mM MgCl2
1M Tris-HCl pH8	10 ml	200mM Tris-HCl
HPLC H2O	<u>35 ml</u>	
Total	<u>50 ml</u>	

Reaction Mix:

10 X buffer	2.5 ul
Exonuclease (20000 U/ml)	0.15 ul
Antartic Phosphatase (25000 U/ml)	0.15 ul
HPLC H2O	<u>7.2 ul</u>
	<u>10.0 ul</u>

Add 10ul of the reaction mix to the DNA on ice. Maintain the plate on ice up to the point that you place it on the thermocycler.

Cycling protocol

96 well and 384 well Cycling (Tetrad and G-Storm):

Lid temp set to 80°C constant

37°C 1 h 80°C 15min 4°C hold

After thermocycle make 1 in 5 dilutions and use diluted DNA for Sequencing.