Kinasing Oligos:

- 1. Take 300 pm of oligo
- 2. Add 5 µl of T4 DNA ligation buffer from NEBL
- 3. Make up the volume to 49 μ l using DD-water
- 4. Add 1 µl of T4 PNK from NEBL.
- 5. Incubate at 37° C for 1 60 min.
- 6. Incubate at 65° C for 20 min to inactivate the enzyme
- 7. Pass it through Bio-Rad spin column to remove unincorporated nucleotides.