Quantify the RNA

- 1. Prepare a blank in micro centrifuge tube (498 μl DNAse/RNAse free water + 2 μl elution buffer
- 2. Sample (498 µl water + 2 µl RNA (premixed)
- 3. Vortex sample

-Leftover RNA add 20 µl ammonium acetate, 200 µl EtOH 100%, store at -20°

Take reading

BioRad Smart Spec 3000

- 1. Press DNA/RNA button :enter
- 2. yes: enter
- 3. turn on vacuum and rinse cuvette with distilled water
- 4. add 500 µl of blank in to cuvette and take blank reading {press read blank}
- 5. empty contents in to vacuum and rinse
- 6. add 500 μl of sample into cuvette and take a reading {press read sample}
- 7. empty contents into vacuum and rinse
- 8. print : 3 full report : exit assay : turn off
- 9. calculation: in excel

RNA concentration $= A_{260} \times 40x$ dilution factor

Purity = A260/A280, Purity should be between 1.8-2.0