

NP1XXXXXN – Buffer Preparation Guide

- 1. Cell Suspension Buffer – 250 mL**
 - a. 12.5mL 1M Tris HCL pH 7.4
 - b. 5.0mL 0.5M EDTA pH 8.0
 - c. Up to 250mL with autoclaved deionized water
 - d. Add 1.5mL enzyme provided then store at 4C

- 2. RNaseA – 2 x 2.8 mL**
 - a. Provided with order

- 3. Cell Lysis Buffer – 250 mL**
 - a. 50mL 1N NaOH
 - b. 12.5mL 20% SDS solution w/v
 - c. Up to 250mL with autoclaved deionized water

- 4. Neutralization Buffer – 250 mL**
 - a. 74g KOAc - Potassium Acetate (CAS no.: 127-08-2)
 - b. Up to 175mL with autoclaved deionized water
 - c. Add 25mL HAC - Glacial Acidic Acid (CAS no.: 64-19-70) to solution
 - d. Using pH meter, pH solution to 5.5 by slowly adding acetic acid
 - e. Top off to 250mL with autoclaved deionized water and store at 4C

- 5. Column Equilibration Buffer – 800 mL**
 - a. 120mL 5M NaCl
 - b. 40mL 1M MOPS pH 7.0
 - c. 120mL 2-Propanol
 - d. Up to 800mL with DNase/RNase-free, low endotoxin sterile water
 - e. Add 1.2mL detergent provided to equilibration buffer

- 6. Wash Buffer – 1.5 liter**
 - a. 300mL 5M NaCl
 - b. 75mL 1M MOPS pH 7.0
 - c. 225mL 2-Propanol
 - d. Up to 1.5L with DNase/RNase-free, low endotoxin sterile water

7. Elution Buffer – 400 mL

- a. 128mL 5M NaCl
- b. 20mL MOPS pH 7.0
- c. 60mL 2-Propanol
- d. Up to 400mL with DNase/RNase-free, low endotoxin sterile water

8. TE Buffer – 30 mL

- a. 300uL 1M Tris HCL pH 7.4
- b. 60uL 0.5M EDTA pH 8.0
- c. Up to 30mL with DNase/RNase-free, low endotoxin sterile water