

## **GTSE1-GFP purification protocol v. 19.03.10**

(adapted from Xmap215 protocol)

### **Cells**

Infect 500mL of SF+ cells at  $1 \times 10^6$ /mL with 200uL BIIC stock (1:2,500 dilution)  
Harvest at peak expression (72 hours)

### **Harvesting Cells**

Spin down SF+ cells for 15min at 1700rpm.  
Resuspend in 40mL Lysis buffer with 1x Pi  
Freeze in 2x~25mL in Falcon tubes.  
Store at -80°C

### **Purification**

Purification is suitable for 50mL of cell suspension. Scale up may require larger columns.

### **Lysis and clarification**

1. Turn on Beckman Ultra Max and set chamber to 4 degrees.
2. Thaw suspension in RT water and transfer to ice
3. Adjust to 10mM CaCl<sub>2</sub>, 1x Pi's
4. Dounce for ten strokes with a pre-chilled dounce
5. Spin for 45' at 80,000 rpm in MLA80 rotor and collect supernatant (tubes fit about 6-7mL each – 8 tubes fit one rotor)
6. Collect supernatant.

### **Nickel column**

7. Add imidazole to 9mM final
8. Load supernatant over pre-equilibrated 5mL His-Trap Nickel column (3% buffer B)
9. Wash with 5 CV of 3% buffer B
10. Wash with 5 CV of high salt buffer (to reduce anion exchange effects)
11. Wash with 5 CV of 10% buffer B
12. Elute with 100% buffer B
13. Run SDS-PAGE to determine peak fractions

### **Gel Filtration Column**

14. Collect peak fractions and pool. Load onto equilibrated Superdex 200 16/60
15. Determine peak fractions by denaturing A280 on NanoDrop.

16. Determine concentration using extinction coefficient: 52,060

17. Adjust to 10% Glycerol, 1mM DTT.

### **Column Set-up**

#### **Nickel Column**

1. Wash out 20% ethanol with 10 CV water
2. Strip column if necessary
  - 10 CV of 50mM EDTA, pH 8.0
  - 8 CV water
  - 1 CV of 100mM NiCl<sub>2</sub>
  - 8 CV water
3. Equilibrate with 10 CV of 3% Buffer B

#### **Gel filtration column**

1. Wash out 20% ethanol with 2 CV water
2. Equilibrate with 2 CV Anion buffer, 100mM KCl

### **Buffers**

#### **Lysis Buffer**

50mM HEPES pH 7.5  
5% glycerol  
0.1% Triton X-100  
200mM NaCl

#### **Ni column buffers**

Buffer A:

25mM Tris-HCl pH 8.0 (3.03g for 1L)  
300mM NaCl (17.53g for 1L)  
20% glycerol

Buffer B:

As above, but with 300mM imidazole (20.4g for 1L)

High Salt Wash

1.5mL Buffer B  
48.5mL Buffer A  
3.0g NaCl

#### **Gel Filtration Buffers**

100mM Anion Buffer

50mM Tris Base(6.1g/L)  
50mM Bis-Tris (14.1g/L)

Adjust to pH 6.6 with HCl

Gel filtration buffer

20mM anion buffer pH6.6

300mM KCl