

BEADCOATING PROTOCOL

- 10^7 particles
- 1 ml 0.1 N MES-Buffer (pH5)
- 2.5 mg EDC (1-Ethyl-3-(3-dimethylaminopropyl)carbodiimid)
- 50 μ g protein (e.g. fibronectin)
- 1% BSA for blocking (100 mg in 10ml PBS)

Protocol:

Wash 10^7 beads in MES-Buffer.

Centrifuge and remove the supernatant carefully.

Resuspend the beads in 500 μ l MES-Buffer and add 12.5 μ l of the EDC solution.

Vortex it and keep it cold on ice.

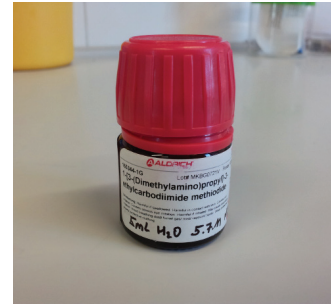
Mix 50 μ g protein with 500 μ l MES-Buffer (cold).

Carefully pipette the protein mixture in the bead solution and vortex it afterwards.

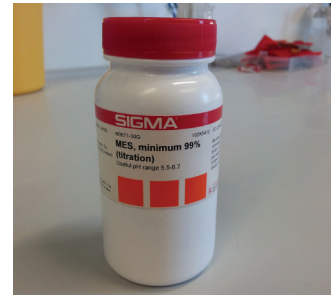
Rotate at 4° and a mix of 1400 over night or longer.

Centrifuge and remove the supernatant carefully.

Add 1ml of 1% BSA solution and store at 4 °C.



EDC (1-Ethyl-3-(3-dimethylaminopropyl)carbodiimid)
Stock: 0.2 g/ml



MES-Buffer