

## *Coating Procedure of Coverslips*

June 11, 2014

### ***Materials***

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- Acid washed coverslips
- Growth Factor Reduced MATRIGEL (BD, 356230)
- 0.01% poly-L-lysine solution (Sigma, P4832)

### ***Procedure***

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#### ***Coating procedure of coverslips with Growth Factor Reduced MATRIGEL Matrix***

1. Thaw Growth Factor Reduced MATRIGEL Matrix. Mix it to homogeneity.
2. Dilute 2  $\mu$ l of MATRIGEL with 98  $\mu$ l of ice-cold PBS or ice-cold medium (total 100  $\mu$ l).
3. Add diluted MATRIGEL to vessel to be coated. Quantity should be sufficient to cover entire growth surface easily (3 ml per 10-cm dish; 1 ml per 6-cm dish; 0.5 ml per well of 6-well plate; 0.2 ml per well of 12-well plate).
4. Incubate at room temperature or 37°C for one hour.
5. **Optional:** Aspirate unbound material gently and dry up the MATRIGEL. Irradiate the MATRIGEL-coated dishes with UV for 15 min.
6. The MATRIGEL-coated dishes are stable for a week when stored at 4°C.

#### ***Coating procedure of coverslips with poly-L-lysine***

1. Incubate acid washed coverslips in 0.01% poly-L-lysine solution for 5 min at room temperature.
2. Remove the coating solution and immediately rinse substrate three times with PBS or serum-free growth medium.
3. Seed cells onto the coated substrate or allow it to dry for use at a later time.