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Oct 11, 2021

Coverslip Functionalization SOP003.v2.2 V.2

Rory Kruithoff¹, Douglas Shepherd¹¹Arizona State University

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protocol .

Human Cell Atlas Method Development Community



Rory Kruithoff
Arizona State University

Document Summary:

This document, SOP003 – Coverslip Functionalization, describes the process for cleaning and modifying the coverslip surface for improved adhesion of a sample. This procedure describes two methods of functionalization, silanization and poly-d-lysine (PDL) coating. Silanization is a method used to promote covalent adhesion of polyacrylamide gel-embedded tissue while poly-d-lysine (PDL) coating of silanized coverslips provides a protein anchoring system which improves adhesion of living cells to the surface of the coverslip.

Quick Overview:

*Part 1 - Clean coverslips**Part 2 - Silanize coverslips**Part 3 - PDL coat coverslips*

Coverslip
Functionalization
SOP003.v2.2.pdf

Rory Kruithoff, Douglas Shepherd 2021. Coverslip Functionalization
SOP003.v2.2. [protocols.io](https://protocols.io/view/coverslip-functionalization-sop003-v2-2-byxxpxpn)
<https://protocols.io/view/coverslip-functionalization-sop003-v2-2-byxxpxpn>
Rory Kruithoff



Coverslip, silanization, poly-d-lysine, coverslip cleaning, tissue adhesion, cell adhesion, allyltrichlorosilane

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Oct 11, 2021

Oct 11, 2021

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v2.2 revision notes

-Added Bottiger lab and OsmFISH references.

References:

Moffitt, J. R., Hao, J., Bambach-Mukku, D., Lu, T., Dulac, C., & Zhuang, X. (2016). High-performance multiplexed fluorescence in situ hybridization in culture and tissue with matrix imprinting and clearing. *Proceedings of the National Academy of Sciences*, 113(50), 14456-14461. <https://doi.org/10.1073/pnas.1617699113>

Boettiger Lab Github:

<https://github.com/BoettigerLab/protocols/blob/master/MatrixEmbedding.md>

OsmFISH [dx.doi.org/10.17504/protocols.io.psednbe](https://doi.org/10.17504/protocols.io.psednbe)

Solution Preparation:

Silanization Solution

- **[M]0.1 % (v/v) triethylamine** (Millipore, TX1200)
- **[M]0.2 % (v/v) allyltrimchlorosilane** (
[Allyltrimchlorosilane Sigma](#)
[Aldrich Catalog #107778](#))
[Chloroform Sigma](#)
- dissolved in chloroform ([Aldrich Catalog #C2432-1L](#))

PDL (Poly-D-Lysine) Solution

- **[M]0.1 Mass Percent Poly-D-Lysine** (molecular weight 30,000-70,000 Da;
[Poly-D-lysine hydrobromide \(PDL\) Sigma](#)
[Aldrich Catalog #P7886](#))
- Dissolved in nuclease-free water.

Additional Reagents:

- [Ethanol Fisher](#)
- [Scientific Catalog #AC615090010](#)
- [Chloroform Sigma](#)
- [Aldrich Catalog #C2432-1L](#)
- [Methanol Sigma](#)
- [Aldrich Catalog #34860-1L-R](#)
- [Hydrochloric acid Sigma –](#)
- [Aldrich Catalog #320331-500ML](#)

For hazard information and safety warnings, please refer to the SDS (Safety Data Sheet).



Using 40-mm-diameter #1.5 coverslips

Part 1 - Clean coverslips

30m

30m

1 

Wash for  **00:30:00** via immersion in 1:1 mix of **[M]37 % (v/v) HCl** and methanol at
 **Room temperature** .

2 

Rinse 3x in DI H₂O.

3 Fill beaker with RNase-free water and autoclave coverslips to sterilize.

4 

Rinse 1x in **70 % Ethanol**.

5 Coverslips can be stored in **70 % EtOH** indefinitely.

Part 2 - Silanize coverslips

6 Dry at **60 °C**.

7 Immerse in silanization solution in Fume-hood for **00:30:00** at **Room temperature**.^{30m}

8 

Wash 1x with chloroform.

9 

Wash 1x with ethanol.

10 Bake in a **60 °C** oven for **02:00:00** to dehydrate the silane layer.^{2h}



11 Remove from oven and let coverslips cool.

12 Store in a desiccated chamber for weeks.


Part 3 - PDL coat coverslips (for cell culture on coverslips)


1h

1h

13 Immerse coverslips in PDL Solution for  **01:00:00** at  **Room temperature** .

14 

Rinse coverslips three times with nuclease-free water and then dry at  **Room temperature** .

15 Silanized, PDL-coated coverslips may be stored at  **Room temperature** in a desiccation chamber for weeks.