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Preparation of tissue culture plates for neural rosettes and neural progenitors

 In 1 collection

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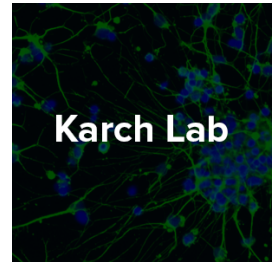
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Protocol status: Working

We use this protocol and it's working

Created: February 18, 2019

Last Modified: February 27, 2019

Protocol Integer ID: 20477



Attachments



IPSC CORTICAL

DIFFER...

179KB

Guidelines

This protocol is part of the IPSC CORTICAL DIFFERENTIATION collection.

This method should be performed using sterile technique.




Materials

Please refer to the attached full manuscript for required materials.

Safety warnings

- ⚠ Please refer to the SDS (Safety Data Sheet) for information about hazards, and to obtain advice on safety precautions.



- 1 Coat 3 wells of a 6-well plate with  1 mL Poly-L-Ornithine (PLO) for  02:00:00 or overnight in humidified incubator.
- 2 Aspirate off PLO and rinse 3 times with DMEM/F12.
- 3 Dilute laminin to 10 $\mu\text{g/mL}$ final in cold DMEM/F12 and add 1 mL/well of solution to PLO coated plates.
- 4 Incubate for  02:00:00 or overnight in humidified incubator.