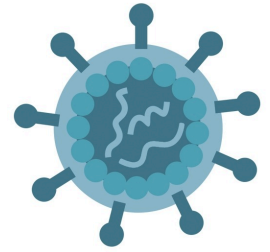


Oct 20, 2019

Viral Preparation

DOI

dx.doi.org/10.17504/protocols.io.8iahuae



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OPEN  ACCESS



DOI: dx.doi.org/10.17504/protocols.io.8iahuae

Protocol Citation: Aditya Mohan 2019. Viral Preparation. **protocols.io** <https://dx.doi.org/10.17504/protocols.io.8iahuae>

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Protocol status: In development

We are still developing and optimizing this protocol

Created: October 20, 2019

Last Modified: October 20, 2019

Protocol Integer ID: 28962






Day 1

- 1 Coat the plates with 1% Poly-L-Lysine solution and leave in dry incubator overnight.


Day 2

- 2 Plate 6E6 HEK293 T cells overnight in DMEM media

Day 3

- 3 In 2 1.5 mL epindorph tube add 150 uL of OptiMEM. In one of them add 15 uL of lipofectamine. In the other add 10 uL of packaging plasmid and 10 uL of the plasmid of interest.
- 4 Incubate for 5 minute.  00:05:00
- 5 Combine the two tubes together for total volume ~ 300 uL and incubate for 5 minutes  00:05:00
- 6 Remove the media from the plate and add the 300 uL dropwise. Add 1 ML of plain DMEM.
- 7 Leave in incubator for 30 minutes  00:30:00
- 8 Add 5 mL of DMEM based media.

Day 5

- 9 Remove the media and add to an eppendorf tube stored in 4 degrees. Add another 5 mL of DMEM based media.  4 °C

Day 7



10 Collect all the media.