

## <u>Standard Operating Procedure</u> for **PICKING iPSC COLONIES**

## 1.0 <u>PURPOSE</u>

This procedure describes how to pick iPSC colonies.

2.0 <u>SCOPE</u> (Should include which Cores this SOP applies to) This procedure applies to all personnel who wish to pick iPSC colonies.

## 3.0 <u>PROCEDURE</u>

- 1. The day before picking ips colonies, prepare gelatin-coated 12-well feeder plates.
- 2. Before picking colonies, add 1 ml hES medium containing 10 µm Y27632 to each well of the plate and incubate it in a 37°C incubator.
- 3. Examine the ips colonies under a microscope in a hood and mark the colonies on the bottom of the dish.
- 4. Cut one colonies into small pieces with a 200  $\mu$ L pipette.
- 5. Transfer the cut pieces to the 12-well feeder plates with a 200  $\mu$ L pipettes.
- 6. Repeat step 4-5 to pick another colony.
- 7. Incubate the plate containing the picked colonies in a 37°C incubator for 24-48 hours.
- 8. After the colonies attach the plates, replace the medium with fresh hES medium and change it every day.

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