

# Human iPSC-Derived Renal Proximal Tubular Cells



## **Table of Contents**

Product Information	2
Preparation of Reagents	3
Culture of Human iPSC-Derived Renal Proximal Tubular Cells	4



## **Product Information**

Catalog. No.	Product Name	Format	Stock Conc.	Storage on Arrival	Thawing Instructions	Storage Once Thawed
ax2115	Human iPSC- Derived Renal Proximal Tubular Cells	≥1 million cells/vial	N/A	Liquid Nitrogen	N/A	N/A
ax3534-250	Renal Epithelial Cell Culture Medium	250 mL	1x	Store at -20°C for up to 6 months	Store at 4°C for up to 1 month	Thaw at 4°C or room temperature

Additional Reagents						
Product Name	Supplier	Product Code				
Growth Factor Reduced Matrigel™	Corning	356230				
Recombinant human bone morphogenetic protein 2 (BMP2)	Sigma-Aldrich	H4791-10UG				
Recombinant human bone morphogenetic protein 7 (BMP7)	Gibco (Thermo Fisher Scientific)	PHC9544				
Y-27632 2HCI (ROCK inhibitor)	Selleck Chemicals	S1049				

These reagents must be added fresh for each aliquot of medium.

### Preparation of Reagents

#### Growth Factor Reduced (GFR) Matrigel™

- Upon receipt, aliquot and store Growth Factor Reduced (GFR) MatrigeI™ at -20°C, according to manufacturer's protocol.
- Coat tissue culture plates 45 minutes 1 hour before thawing **Human iPSC-Derived Renal Proximal Tubular Cells**.

#### Recombinant Human Bone Morphogenetic Protein 2 (BMP2)

• Prepare a 10 μg/mL stock solution of BMP2 by resuspending 10 μg of the lyophilized powder in 1 mL Dulbecco's-phosphate-buffered saline (D-PBS) with 0.05 % human serum albumin (HSA).

#### Recombinant Human Bone Morphogenetic Protein 7 (BMP7)

• Prepare a 5 μg/mL stock solution of BMP7 by resuspending 10 μg of the lyophilized powder in 2 mL Dulbecco's-phosphate-buffered saline (D-PBS) with 0.05 % HSA.

Differentiation Factor	Stock Concentration	Final Concentration	In 50 mL Medium
BMP2	10 μg/mL	10 ng/mL	50 μL
BMP7	5 μg/mL	2.5 ng/mL	25 μL

#### **Renal Epithelial Cell Culture Medium**

- Upon receipt, aliquot and store Renal Epithelial Cell Culture Medium at or below -20°C protected from light.
- When ready to use, thaw an aliquot of Renal Epithelial Cell Culture Medium overnight at 4°C or room temperature
  in the dark.



## Culture of Human iPSC-Derived Renal Proximal Tubular Cells

#### Coating

- Thaw aliquots (as needed) of GFR MatrigeI™ on ice before use.
- Dilute **GFR MatrigeI™** 1:50 in ice-cold serum-free medium (DMEM or another suitable medium) on ice to make a 1x working solution e.g. 100 μL of **GFR MatrigeI™** into 5 mL of serum-free medium.
- Coat the surface of your culture vessel with the GFR Matrigel™ 1x working solution. We recommend coating at a volume of 150 µL per cm².
- Incubate the coated cell culture vessel at 37°C, 5% CO, in a humidified incubator for 1 hour.

Consult with manufacturer's protocol for further detailed instructions on coating plates with **GFR Matrigel™**.

#### **Thawing and Plating**

- Prepare a sufficient volume (dependent on the culture vessel format for plating) of Renal Epithelial Cell Culture
   Medium supplemented with 10 μM Y-27632 2HCl and warm to 37°C prior to use.
- Before thawing the cells prepare a biological safety cabinet, spraying the vial and hood thoroughly with 70% ethanol
  and wiping with an autoclaved paper towel before placing the vial in the hood.
- To thaw the cells transfer the vial of cells from storage by transporting the vial buried in dry ice. Remove the vial from dry ice and transfer it to a 37°C water bath.
- Quickly thaw the vial of cells in a 37°C water bath. Do not completely submerge the vial (only up to 2/3rd of the vial).
   Remove the vial before the last bit of ice has melted, after 1-2 minutes.
- Do not shake the vial during thawing.
- Take the vial of cells to the prepared biological safety cabinet.
- Prepare a 15 mL sterile conical tube with 9 mL of pre-warmed, 37°C, Renal Epithelial Cell Culture Medium + 10μM Y-27632 2HCI.
- Using a P1000 pipette, quickly and gently add the cell suspension drop-wise into the medium. This must be done
  while there is still a small ice lump in the cryovial.
- Gently wash the cryogenic vial with 1 mL of Renal Epithelial Cell Culture Medium + 10 μM Y-27632 2HCl to
  ensure all of the cells are transferred to the 15 mL sterile conical tube.
- Centrifuge the cells at **300** x *g* for **5** minutes at room temperature.
- Carefully remove the supernatant and resuspend the cell pellet in 1 mL of pre-warmed, 37°C, Renal Epithelial Cell Culture Medium + 10 μM Y-27632 2HCI.
- Perform a cell count to determine the number of viable cells and ensure optimal seeding density.
- Dilute the cells into the required volume of pre-warmed, 37°C, Renal Epithelial Cell Culture Medium + 10 μM
   Y-27632 2HCI.
- Seed cells into the culture vessel at the recommended seeding density of 50,000 viable cells/cm<sup>2</sup>. The day of seeding the cells is Day 0.
- Incubate the cells at 37°C, 5% CO, in a humidified incubator for 2 days.

#### **Maintenance**

- On Day 2 (48 hours after seeding the cells), the culture medium should be replaced with Renal Epithelial Cell
   Culture Medium containing BMP growth factors (BMP2 and BMP7) without Y-27632 2HCI.
- Supplement the required volume of Renal Epithelial Cell Culture Medium with 10 ng/mL BMP2 and 2.5 ng/mL BMP7. Warm to 37°C prior to use.
- Remove the spent cell culture medium from the culture vessel and replace with pre-warmed, 37°C, Renal Epithelial
   Cell Culture Medium + 10 ng/mL BMP2 + 2.5 ng/mL BMP7.
- On Day 4 (96 hours after seeding the cells), the cells will be ready to use for endpoint assays.

Got any questions? Need help with the protocol?

Contact Axol Technical Support at <a href="mailto:support@axolbio.com">support@axolbio.com</a>

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