

## HEK293-PSMA Cells | 305992

### General information

#### Description

**Disclaimer: The prices displayed for cell lines are exclusively for academic/not-for-profit customers. For commercial entities the price is approximately €6,250.**

**If you represent a commercial entity or are unsure which category applies, please [contact us](#).**

HEK293-PSMA cells are human embryonic kidney 293 (HEK293) cells engineered to stably express human prostate-specific membrane antigen (PSMA), also known as glutamate carboxypeptidase II (FOLH1/GCPII). PSMA is a type II transmembrane glycoprotein with enzymatic folate hydrolase and carboxypeptidase activity that is highly expressed in prostate cancer, particularly in advanced, metastatic, and castration-resistant disease. In addition to prostate malignancies, PSMA expression has also been observed in the neovasculature of various solid tumors. Because of its strong tumor-associated expression and accessible extracellular domain, PSMA has become a major target for diagnostic imaging, radioligand therapy, antibody-based therapeutics, and engineered immune cell approaches.

HEK293-PSMA cells are widely used in oncology research and therapeutic development for characterization of PSMA-targeted monoclonal antibodies, antibody-drug conjugates, radiopharmaceuticals, bispecific T-cell engagers, CAR-T cell therapies, and small-molecule inhibitors. The stable recombinant expression system enables quantitative analysis of ligand binding, receptor occupancy, antigen density, internalization kinetics, and target-dependent cytotoxicity. These cells are particularly valuable for evaluating PSMA-directed imaging probes and radioligand platforms, as PSMA undergoes efficient internalization following ligand engagement. Additional applications include flow cytometry assay development, uptake studies, reporter assays, high-throughput screening, and validation of targeted delivery systems for prostate cancer therapeutics.

**Organism** Human

**Tissue** Fetal Kidney

### Characteristics

**Age** Fetus

**Gender** Female

**Morphology** Epithelial-like

**Growth properties** Monolayer, adherent

### Regulatory Data

**Citation** HEK293-PSMA (Cytion catalog number 305992)

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**Biosafety level** 1**NCBI\_TaxID** 9606**Biomolecular Data****Receptors expressed** PSMA**Handling****Culture Medium** RPMI 1640, w: 2.0 mM stable Glutamine, w: 2.0 g/L NaHCO<sub>3</sub> (Cytion article number 820700a)**Supplements** Supplement the medium with 10% FBS, 1 mM sodium pyruvate, 10 mM HEPES, 1% NEAA. Add Geneticin (G418-Sulfat) to achieve a final concentration of 1 mg/mL.**Dissociation Reagent** Trypsin-EDTA**Subculturing** For routine adherent cell culture: Aspirate the old culture medium from the adherent cells, and wash them with PBS to remove any remaining medium. After aspirating the PBS, add the appropriate volume of Trypsin/EDTA solution based on the culture vessel size (e.g., 1 ml for a T25 flask, 3 ml for a T75 flask) and incubate at room temperature or 37°C until the cells detach (5-10 minutes). Monitor detachment under a microscope, and gently tap the vessel if necessary to release the cells. Once detached, add complete medium to inactivate the Trypsin/EDTA, gently resuspend the cells, and transfer an aliquot of the cell suspension into a new culture vessel containing fresh medium. Place the vessel in an incubator set to 37°C with 5% CO<sub>2</sub>, and change the medium every 2-3 days.**Fluid renewal** 2 to 3 times per week**Post-Thaw Recovery** After thawing, split the cells at a ratio of 1:2 to 1:3 in T25 flasks and allow the cells to recover from the freezing process and to adhere for at least 24 hours.  
  
For best attachment and viability after thawing the cells, we recommend using Collagen-coated flasks or plates for the initial seeding after cryo-recovery. Collagen coating is not required for subsequent routine culture of the cells.**Freeze medium** As a cryopreservation medium, we use complete growth medium (including FBS) + 10% DMSO for adequate post-thaw viability, or CM-1 (Cytion catalog number 800100), which includes optimized osmoprotectants and metabolic stabilizers to enhance recovery and reduce cryo-induced stress.

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### Thawing and Culturing Cells

1. Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit.
2. Upon receipt, either store the cryovial immediately at temperatures below  $-150^{\circ}\text{C}$  to ensure the preservation of cellular integrity, or proceed to step 3 if immediate culturing is required.
3. For immediate culturing, swiftly thaw the vial by immersing it in a  $37^{\circ}\text{C}$  water bath with clean water and an antimicrobial agent, agitating gently for 40-60 seconds until a small ice clump remains.
4. Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening.
5. Carefully open the disinfected vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of room-temperature culture medium, mixing gently.
6. Centrifuge the mixture at  $300 \times g$  for 3 minutes to separate the cells and carefully discard the supernatant containing residual freezing medium.
7. Gently resuspend the cell pellet in 10 ml of fresh culture medium. For adherent cells, divide the suspension between two T25 culture flasks; for suspension cultures, transfer all the medium into one T25 flask to promote effective cell interaction and growth.
8. Adhere to established subculture protocols for continued growth and maintenance of the cell line, ensuring reliable experimental outcomes.

### Incubation Atmosphere

$37^{\circ}\text{C}$ , 5%  $\text{CO}_2$ , humidified atmosphere.

### Shipping Conditions

Cryopreserved cell lines are shipped on dry ice in validated, insulated packaging with sufficient refrigerant to maintain approximately  $-78^{\circ}\text{C}$  throughout transit. On receipt, inspect the container immediately and transfer vials without delay to appropriate storage.

### Storage Conditions

For long-term preservation, place vials in vapor-phase liquid nitrogen at about  $-150$  to  $-196^{\circ}\text{C}$ . Storage at  $-80^{\circ}\text{C}$  is acceptable only as a short interim step before transfer to liquid nitrogen.

## Quality Control & Molecular Analysis

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### **Sterility**

Mycoplasma contamination is excluded using both PCR-based assays and luminescence-based mycoplasma detection methods.

To ensure there is no bacterial, fungal, or yeast contamination, cell cultures are subjected to daily visual inspections.