Glatte muskelceller fra menneskelig lungearterie (HPASMC)

300706

GENERELL INFORMASJON

Beskrivelse

Primary Human Pulmonary Artery Smooth Muscle Cells (HPASMC) are isolated from the pulmonary artery. They are a key model for pulmonary arterial hypertension, including hypoxia-driven proliferation and vascular remodeling.
Hypoxia drives the proliferation of HPASMC, and PDGF-BB shifts them toward a synthetic, migratory phenotype, reproducing key features of pulmonary vascular remodeling. Because the balance of BMP and TGF-β signaling is central to pulmonary arterial hypertension, the cells are also used to study BMPR2-related mechanisms.

Vev	Lung (pulmonary artery
Sykdom	Normal pulmonary arterial smooth muscle (non-tumorigenic primary cell)
Metastaseområde	Not applicable (normal, non-tumorigenic primary cell)

Søknader

Pulmonary arterial hypertension: BMP/TGF-β balance, proliferation and migration; Hypoxia-induced proliferation, PPARγ and microRNA-21; Long non-coding RNA regulation of migration in pulmonary arterial hypertension; Functional testing of BMPR2 promoter variants in reporter assays; PDGF-BB-induced phenotype switching and metabolomics; Endothelin-1 signaling and proteomics of BMPR2-mutant cells

Synonymer

- HPASMC
- Human pulmonary artery smooth muscle cells
- provitro 1210613

DONOR

Donorens alder	Lot-specific (see CoA)
Etnisitet	Lot-specific (see CoA)

KLASSIFISERING

Organisme	Menneske
Sex	Tomtespesifikk
Vekstegenskaper	Tilhenger
GMO-status	Uten GMO

Detaljer om GMO-status

No genetic modification; primary human pulmonary artery smooth muscle cells isolated from normal pulmonary arteries. Not transformed or immortalized.

Biosikkerhetsnivå	1

EGENSKAPER

Morfologi	Spindle-shaped (smooth muscle)
Celletype	Primary smooth muscle cells

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Tumorinduserende	No
Fordoblingstid	Not specified by AddexBio; typical primary VSMC doubling time 36–72 hours

HÅNDTERING OG KULTUR

Næringsmedium

Smooth Muscle Cell Growth Medium kit (Cytion article number 850075); Smooth Muscle Cell Growth Medium mix (Cytion article number 850074)

Inkubasjonsatmosfære	37°C, 5% CO_2, saturated humidity.

Kosttilskudd

Supplement as specified in the AddexBio SMC supplement kit (refer to specification sheet). For cryopreservation: complete culture medium + 5% (v/v) DMSO.

Dissosiasjonsmiddel	Subculture reagents as recommended on the certificate of analysis

Subkultivering

Subculture the cells at about 75 % confluence. Denser cultures undergo irreversible contact inhibition and stop proliferating. Use the subculture reagents recommended on the certificate of analysis.

Oppdelingsforhold	1 to 3
Såtetthet	More than 4,000 cells/cm²
Væskefornyelse	24 hours after seeding, then every 48 to 72 hours
Gjenoppretting etter tining	Post-thaw viability ≥ 70 % viable cells. More than 80 % confluence within the expected recovery period.

Tining og dyrking

Thaw the cryovial quickly in a 37 °C water bath, swirling gently. Just before the last ice has melted, move the vial to a laminar flow hood and decontaminate its outer surface.
Transfer the cell suspension into a 15 ml centrifuge tube, add 10 ml of Smooth Muscle Cell Growth Medium and centrifuge at 250 × g for 5 minutes.
Remove the supernatant and resuspend the cell pellet in 10 ml of Smooth Muscle Cell Growth Medium. Check cell number and viability.
Divide the suspension between two 25 cm² culture flasks (5 ml each). Using two flasks reduces the risk of losing all cells if a passage fails.
Incubate at 37 °C, 5 % CO_2 and saturated humidity, with the screw caps opened by half a turn for gas exchange.
Change the medium after 24 hours to remove unattached cells, then every 48 to 72 hours, using pre-warmed medium only.
Subculture the cells at about 75 % confluence. Denser cultures undergo irreversible contact inhibition and stop proliferating. Use the subculture reagents recommended on the certificate of analysis.
Recommended seeding density: more than 4,000 cells/cm².

KVALITETSKONTROLL OG KRYOKONSERVERING

Sterilitet

Mycoplasma contamination is excluded by PCR.
To ensure there is no bacterial, fungal or yeast contamination, cultures are subjected to visual inspection.

Frysemedium	As a cryopreservation medium, we use complete growth medium + 10% DMSO for adequate post-thaw viability.

Oppbevaringsbetingelser

For long-term preservation, store vials in liquid nitrogen at about −150 to −196 °C. Storage at −80 °C is acceptable for up to one week only.

Fraktvilkår

Cryopreserved primary cells are shipped on dry ice. On receipt, check the cryovial for damage and transfer it without delay to appropriate storage.

UTTRYKK OG MOTTAKELIGHET

Proteinekspresjon

Smooth muscle α-actin; calponin; smooth muscle myosin heavy chain

Virus

Free of HIV, Hepatitis B, Hepatitis C (confirmed negative by AddexBio)

IDENTIFIKATORER

Kildehenvisning

Human pulmonary artery smooth muscle cells (HPASMC) (Cytion catalog number 300706)

NCBI-skattenummer	9606
Cellosaurus	Not assigned

JURIDISK OG SIKKERHET

Sikkerhetsregler

When planning to store a cryovial in liquid nitrogen for future thawing, it is mandatory to adhere to stringent safety measures. Appropriate protective gloves and clothing are essential, and the use of a face mask or safety goggles is required during the transfer of frozen samples to or from the liquid nitrogen tank. This is to mitigate the risk of injury from potential cryovial explosions upon removal, which can result in the projection of sharp fragments. Primary cells are of human origin, and no known test procedure can ensure the complete absence of infectious agents. Handle all products as potentially infectious.

Garanti

At Cytion, we stand behind the quality and viability of our products.
We guarantee that our cells will be viable at the start of culture for
60 days after shipment from our facilities.
We assure that all materials conform to the specifications described in the
Certificate of Analysis and/or Product Sheet.
For full details, please refer to our
Warranty Statement.

Avtale om materialoverføring kreves	Yes

PRODUKTDETALJER

Sammensetningstabell