SCINUS CELL EXPANSION

CONTROLLING GROWTH

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SCINUS CELL EXPANSION

OUR VISION: MAKING CELL THERAPY AND REGENERATIVE MEDICINE ACCESSIBLE AND AVAILABLE TO EVERYONE, STRIVING TO POSITIVELY CHANGE HEALTH CARE.

OUR MISSION: TO PROVIDE TECHNOLOGY AND PROCESSES TO ENABLE AND IMPROVE CELL THERAPY AND REGENERATIVE MEDICINE.

Controlable upscaling, affordable costs





Controlable upscaling, affordable costs



CELL EXPANSION TECHNOLOGY



Controlable upscaling, affordable costs





Controlable upscaling, affordable costs



PROCESS DEVELOPMENT



Controlable upscaling, affordable costs



CELL EXPANSION TECHNOLOGY





Unique, volume expandable technology

Continuous perfusion, Extremely flexible



Unique, volume expandable technology



100 mL





1400 mL







Continuous perfusion, Extremely flexible







Easy to use GUI



Unique, volume expandable technology

Continuous perfusion, Extremely flexible



Easy to use GUI







Single-use culture bag

one SCINUS cabinet for ADHERENT as well as SUSPENSION cell types





Dynamic culture

Homogenous and low shear environment; Customizable



PARAMETER	RANGE
Rocking angle	+/- 100 °
Rocking velocity	0 - 500 °/s
Static interval	0 to 100 hours















- Direct culture of bone marrow-derived stromal cells (BMSCs)
- Direct culture of adipose-derived stem cells (ASCs)
- Muscle Progenitor Cells, UC, WJ, Periosteum derived cells
- Exosome / EV production

















- T-Cell cultivation for CAR-t
- CD34⁺ cell culture



Donor 2 - T flask Donor 2 - Scinus

Donor 1 - T flask Donor 1 - Scinus













- Large scale spheroids culture (e.g. iPSC)
- Organoid culture





Controlable upscaling, affordable costs



PROCESS DEVELOPMENT



- Microcarrier process considerations
 - Seeding efficiency
 - Cell motility
 - Minimize aggregation
 - Harvest efficiency
- Microcarrier process parameters
 - Seeding density
 - Agitation regime
 - Expansion timing
 - Microcarrier concentration





- Investigate efficiency in multiple microcarriers
- Investigate attachment kinetics

Microcarrier coating	1h	2h	4h	24h	
Collagen					
Vitronectin					X



- Investigate efficiency in multiple microcarriers
- Investigate attachment kinetics
- Investigate allowable seeding densities





- Use Design-of-Experiments for multi-parameter optimization where appropriate
- Design space for e.g. agitation rate, expansion timing, MC density
- Use quantifiable responses (e.g. PDT, distribution)









Total cell number

- Day 3 expansion 2 stops/day
- Day 3 expansion 4 stops/day
- Day 3 expansion 6 stops/day
- Day 5 expansion 2 stops/day
- Day 5 expansion 4 stops/day
- Day 5 expansion 6 stops/day
- Day 7 expansion 2 stops/day
- ▲ Day 7 expansion 4 stops/day
- -▼ Day 7 expansion 6 stops/day



• Day of expansion is not a significant term, agitation regime is





- Optimize for model terms (PDT and distribution)
- Optimum at day 5*, and 4 pauses per 24 hours



* Not a significant factor





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