

Research Paper

Generation of Human 3D Airway Assembloids for Advanced Modeling

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Received: 2025.03.18; Accepted: 2025.08.23; Published: 2025.10.01

Abstract

The development of physiologically relevant *in vitro* 3D models is crucial for studying lung biology and disease mechanisms. While airway organoids have significantly improved our ability to mimic lung tissue, they lack key nonepithelial components that are essential for tissue homeostasis. Here, we describe the generation of human airway assembloids, combining airway organoids, stromal fibroblasts, and endothelial cells to better replicate the native lung environment. The model was generated from healthy lung tissue donors by using a scaffold-free culture system to promote cell self-organization. Assembloids exhibited long-term viability, maintained typical airway epithelial markers, and demonstrated functional characteristics, such as mucus production and ciliary beating. This technology provides a powerful platform for studying airway physiology, disease mechanisms, and therapeutic approaches, with potential applications in regenerative and personalized medicine. Our study established a novel, reproducible 3D assembloid model of the human airways, bridging the gap between traditional organoid cultures and complex tissue engineering strategies.

Keywords: lung, airway, 3D model, organoids, assembloids

Introduction

Organ functionality is dependent on synergistic interactions between multiple cell types within a three-dimensional (3D) structure [1]. Most *in vitro* cell culture techniques still rely on two-dimensional (2D) monolayers although the transfer of cells from their native environments to 2D cultures leads to losses of tissue-specific cellular functions and architectural cues [2,3]. Consequently, resembling a 3D architecture is essential for developing more accurate *in vitro* model systems [4,5].

In the last decade, our understanding of human stem cell biology has grown and the availability of 3D cell culture technologies has increased significantly, enabling the development of stem cell-derived organoid cultures that mimic organ structure and function. Organoids are typically derived from either pluripotent or tissue-specific adult stem cells (e.g., from the brain [6], liver [7], pancreas [8,9], and lung [10,11]) that are cultured *in vitro* under specific conditions to promote self-organization similar to the

architectures of the organs they represent. Therefore, organoids are innovative 3D models that recapitulate key features of organ physiology, including intercellular interactions, tissue organization and functionality, thus facilitating studies of organ development and pathogenesis, drug discovery and regenerative medicine [12–16].

3D cell culture systems provide significant advantages over traditional 2D cultures because they replicate native tissue architecture more reliably. Although organoids represent a major advance in the 3D culture field, they remain simplified models of natural tissues. Notably, they lack key nonepithelial components, and particularly fail to recapitulate the cell-cell interactions of the natural organ. Functional networks between epithelia and surrounding tissues are essential in the regulation of development and in organ-specific pathogenesis. Replication of the natural microenvironment that includes fibroblasts and endothelial cells is expected to generate high-fidelity models that better mimic the natural organ.

Assembloids have emerged as next-generation 3D tissue models that integrate multiple cell types and offer a higher fidelity representation of organ complexity and function. First introduced by Pasca [17], the term “assembloid” refers to a 3D multicellular model that combines multiple organoids or specialized cell types. These advanced models structurally mimic key physiological and pathological processes such as cell migration and intercellular signaling.

Assembloids that display liver, kidney, heart, and brain tissues are available [18]. However, to our knowledge, no assembloids of healthy pulmonary tissue have been created, although lung cancer assembloids have already been described [19]. Herein, we report the development of a functional assembloid that recapitulates the healthy airways by combining an airway organoid with a stromal fibroblast compartment, isolated from the same tissues, and a human endothelial cell line. This cellular composition enabled a more accurate representation of the native airway microenvironment that includes intercellular interactions, essential to organ homeostasis.

Materials & Methods

Sample collection

Healthy human lung tissue was obtained from the Thoracic Surgery and Lung Transplant Unit, Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, Milan, Italy. The use of human specimens was approved by the Ethical Committee of Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano (CE-0006851). Young donors

underwent minimal invasive wedge lung resection indicated for spontaneous pneumothorax.

Human airway organoid generation and culture

Human airway organoids (hAO) were generated from tissue-resident adult stem cells (Fig. Supplementary 1A) as previously described [11]. After 10–21 days of culture, hAO were removed from the Cultrex-UltiMatrix RGF-BME (BME001-05, Bio-Techne) in cold Dulbecco's phosphate buffered saline without Ca^{2+} and Mg^{2+} (DPBS, ECB4004L Euro Clone) and enzymatically dissociated with trypsin-EDTA solution (10X, T4174 Sigma-Aldrich) for 5 min. Cells were then counted in a Burkert chamber and seeded in Cultrex-UltiMatrix RGF-BME (ratio 1:6-1:12), enabling the formation of new organoids. Unless specified, hAO were analyzed after at least 15–20 days post-seeding at the indicated passage. hAO were characterized by RT-qPCR, flow cytometry, immunohistochemistry, and immunofluorescence staining at different passages. All cell cultures were routinely tested for mycoplasma contamination by MycoAlert Assay Control Set (LT07-518, Lonza).

Lung fibroblast isolation and culture

Lung fibroblasts (LF) were isolated from the same lung biopsy specimens. Once minced, one or two pieces were seeded on 6-well culture plate with high-glucose Dulbecco's Modified Eagle Medium (DMEM, ECM0102L, Euro Clone) supplemented with 1% GlutaMAX (35050-038, Gibco), 10% fetal bovine serum (FBS, 10270-106, Gibco), and 1% antibiotic-antimycotic solution (ECM0010D, Euro Clone). After 6–8 days, fibroblasts began migrating from the tissue fragments and adhered to the plastic surface, establishing a classical stromal monolayer (Fig. Supplementary 1B). LF were routinely grown on DMEM 10%FBS, 1%GlutaMAX, 1% penicillin-streptomycin (P4333, Sigma-Aldrich) at 37°C in a humidified atmosphere of 5% CO_2 and passaged with 0,05% trypsin (25-054-CI, Corning) and 0,02% EDTA (E7889, Sigma-Aldrich). RT-qPCR, flow cytometry, and immunofluorescence were used to confirm their identity. Cultured fibroblasts (80% confluence) were used for experiments from passages 3 to 6.

RT-qPCR analysis

Total RNA was isolated from at least five independent hAO using TRIzol reagent (FS-881, Fisher Molecular Reagent) according to the manufacturer's instructions. RNA concentration was measured using a NanoDrop ND-100

spectrophotometer (NanoDrop Technologies). To perform an RT-qPCR assay, cDNA was synthesized from 500ng of total RNA with SuperScript IV VILO Master Mix with ezDNase (11766050, Invitrogen). The cDNA was diluted 10-fold, and 1µL of the sample was used as a template for RT-qPCR analysis with PowerUp SYBR Green Master Mix (A25742, Applied Biosystems) on a QuantStudio3 Real-Time PCR Instrument (A28131, Applied Biosystems). Relative expression levels of the specific target genes (Table 1) [20,21] were determined using the $2^{-\Delta\Delta C_t}$ method and normalized to the geometric mean of the GAPDH and HPRT mRNA levels.

Flow cytometry

hAO were dissociated with trypsin-EDTA solution 1X in DPBS for 5 min to obtain a single-cell suspension. A minimum of 100,000 cells per tube were stained for each sample. For surface marker analysis, cells were incubated with fluorophore-conjugated antibodies in a total volume of 100µL of DPBS for 10 min in the dark at room temperature (RT), washed with 1ml of DPBS, and resuspended in 200µL of PBS. For intracellular marker analysis, cells were fixed in 500µL of 1% paraformaldehyde (PFA,15710, ChemCruz) for 15 min at RT, washed with 2ml of DPBS, permeabilized and incubated with 500µL of 0.05% saponin (SAE0073-10G, Sigma-Aldrich), 2% FBS, and the primary antibody. Next, permeabilized and stained cells were washed with 1ml of 0.05% saponin, 2% FBS, and incubated with the secondary antibody for 10 min at RT. After two additional washing steps, cells were resuspended in 200µL of DPBS.

LF were collected, washed once with DPBS, resuspended in 100µl staining solution (DPBS, 1% BSA containing specific antibodies), and kept for 20 min in the dark at 4 °C. Cells were then washed and resuspended in 200µl of DPBS 1% BSA solution. Flow cytometry of LF was completed using the same protocol described above for hAO. Monoclonal antibodies are reported in Table 2.

Stained cell suspensions of hAO and LF were analyzed with the FACSCanto II cytometer with FACSDiva analysis software (Becton Dickinson, BD) or with the BD FACS Lyric (BD) using FlowJo software (BD). Acquired events were plotted against forwarding scatter-height and forward scatter-area (FSC-A) to exclude cell doublets (P1 gate) and then against FSC-A and side scatter-area physical parameters to select live cell populations excluding debris (P2 gate). At least 10,000 P2 events were acquired. Markers of interest were measured in histograms or dot plots by an analytical gate for marker-positive events. Histogram data were

presented as mean $\pm$ SEM of at least n=3 independent experiments.

MTT assay

hAO were washed twice with DPBS without Ca^{2+} and Mg^{2+} and 0.5 mg/mL of thiazolyl blue tetrazolium bromide (M5655; Sigma-Aldrich) dissolved in DMEM without phenol red (D1145; Sigma-Aldrich) (MTT Solution) was added to each well. hAO were then incubated in the dark at 37 °C with 5% CO_2 in a humidified atmosphere for 2 h. Next, the MTT solution was removed and 96% ethanol (528,151; Carlo Erba) was added to each well. Plates were next placed on a rocking shaker at 37 °C with 5% CO_2 in a humidified atmosphere for at least 2 h. Optical density was measured at 570 nm on a microplate reader (TECAN Infinite 200 PRO).

Table 1. Primers

E-Cad (E-Cadherin)	FWD	TGCTCTTCCAGGAACCTCTG
	REV	GCGGCATGTAGGTGTCA
ELF3 (E74 like ETS transcription factor 3)	FWD	CTCTTCCCCAGCGATGGTIT
	REV	CCCTCGAGACAGTCCCAGTA
KRT5 (Keratin 5)	FWD	GCATCACCCTTCTGGGTAA
	REV	GACACACTTGACTGGCGAGA
p63 (Transformation-related protein 63)	FWD	CCAGAGATGGGCAAGTCTCTG
	REV	ACTGTCCGAAACTTGCTGCT
SCGB1A1 (Secretoglobulin Family 1A member 1)	FWD	TCCTCCACCATGAAACTCGC
	REV	AGGAGGGTTTCGATGACACG
AGR2 (Anterior gradient protein 2 homolog)	FWD	TCAGAAGCTTGGACCGCATC
	REV	AGTGTAGGAGAGGGCCACAA
NPHP1 (Nephrocystin-1)	FWD	CAGAGCCACATGGCAACCTA
	REV	ACCCAGCCACAGCTTAACCTC
DNAH5 (Dynein axonemal heavy chain 5)	FWD	AGAGGCCATTGCGAAACG
	REV	CCCGGAAAATGGGCAAACTG
SOX2 (SRY-sex determining region Y-box 2)	FWD	CACATGAAGGAGCACCCGGATTAT
	REV	CGGGAAGCGTGTACTTATCCTTCT
NKX2.1 (NK2 homebox 1)	FWD	ACCAAGCGCATCCAATCTCA
	REV	CAGAGCCATGTCAGCACAGA
LGR5 (Leucine-rich repeat-containing G protein coupled receptor 5)	FWD	AGGCCTTGTCTGGGTGAA
	REV	TGGCTTCACTGGGTACTGTTT
CD73 (Cluster of differentiation 73)	FWD	CACTGGGACATTCGGGTTTT
	REV	CGTCCACACCCCTCACTTTC
CD90 (Cluster of differentiation 90)	FWD	CGAACCAACTTCACCAGCAAAT
	REV	CCTTGCTAGTGAAGCGGATA
CD105 (Cluster of differentiation 105)	FWD	CCGCGCTTCAGCTTCCT
	REV	GAGGGTGCCGGTTTTGG
VIM (Vimentin)	FWD	AGGCAAAGCAGGAGTCCACTGA
	REV	ATCTGGCGTTCAGGGACTCAT
ACTA2 (α Smooth Muscle Actin 2)	FWD	CTGTTCAGCCATCCTTCAT
	REV	TCATGATGCTGTGTAGGTGGT
CFTR (Cystic Fibrosis Transmembrane Conductance Regulator)	FWD	CACAGGACAGCCCTTCTTTC
	REV	TGCCCATGGCTATCTACTT
ENaC - α (Epithelial Sodium Channel - α)	FWD	GCTGATAACCAGGACAAAACACAA
	REV	CGTCGCTGGGCAGGAA

Table 2. Antibodies

Antibody Target	Host Animal/species	Fluorophore(s)/Conjugate	Clone	Cat. Number	Vendor/ company
anti-EpCAM	Mouse	PECy7	1B7	25-9326-42	eBioscience
anti-CD271	Mouse	Alexa 647	C40-1457	560326	BD
anti-CD66c	Mouse	PE	KOR-SA3544	12-0667-42	eBioscience
anti-TSPAN8	Rat	Alexa 405	458811	FAB4734V	R&D
anti-FOXJ1	Mouse	-	2A5	14-9965-82	eBioscience
anti-CD90	Mouse	PECy7	5E10	561558	BD
anti-CD73	Mouse	PE	AD2	550257	BD
anti-CD105	Mouse	PerCP-Cy5.5	266	560819	BD
anti-Vimentin	Mouse	-	VIM 3B4	CBL202-K	Merck-Chemicon
anti-p63	Rabbit	-	EPR5701	ab124762	Abcam
anti-CC10/SCGB1A1	Mouse	-	E-11	sc-365992	Santa Cruz
anti-KRT5	Mouse	-	XM26	MA5-12596	Invitrogen
anti-beta IV tubulin	Rabbit	-	EPR16775	ab179504	Abcam
anti-HT2-280	Mouse	-	N/A	TB27AHT2-280	Terrace Biotech
Anti-MUC5AC	Mouse	-	45M1	sc-21701	Santa Cruz
anti-CFTR	Mouse	-	CF3	NB300-511	Novus Biologicals
Anti-EnaC α	Rabbit	-	Polyclonal	orb101489	Biorbyt
Anti-rabbit	Goat	Alexa 555	-	AB-143157	Invitrogen
Anti-mouse	Goat	Alexa 488	-	Ab-2534069	Invitrogen
Anti-mouse	Goat	Alexa 555	-	A21-422	Invitrogen
IgG Isotype control	Mouse	-	-	31903	Invitrogen

Immunohistochemistry and immunofluorescence staining

hAO were washed with DPBS and fixed for 15-30 min in cold 4% PFA, followed by dehydration, paraffin embedding, sectioning (3 μ m thickness), and standard hematoxylin and eosin (H&E), and Alcian-blue periodic acid Schiff (AB-PAS) staining protocols. Images were acquired using the Leica Aperio AT2 (objectives 20X and 40X).

Unstained 3 μ m sections were deparaffinized and incubated overnight at 4°C with primary antibodies. On the following day, samples were washed and incubated with secondary antibodies and DAPI (1 μ g/ml in DPBS, ThermoFisher Scientific). ProLong Gold Antifade Reagent (P36930, ThermoFisher Scientific) was used as a mounting medium. Images were obtained on a confocal microscope (Leica SP8, 20X objective).

To conduct immunofluorescence staining of hAO, samples were removed from the extracellular matrix (ECM), washed with DPBS, fixed with 4% PFA for 40 min on ice, permeabilized with 0.3% Triton X-100 (9036-19-5, Sigma-Aldrich) in DPBS for 40 min at RT, washed again with DPBS, and incubated three times for 10 min with 100mM glycine (1610724, Bio-Rad) in DPBS at RT. After another wash in DPBS, they were blocked for 12h at RT with blocking solution containing 0.1% BSA (11922.03, Serva), 0.2% Triton X-100, 0.05% Tween-20 (P7949, Sigma Aldrich),

and 10% goat serum (S26, ThermoFisher Scientific) in DPBS.

Samples were incubated with primary antibodies diluted in blocking solution for 24h at 37 °C in a shaker, then washed carefully with DPBS and incubated with secondary antibodies and DAPI for 2 h at 27 °C. Samples were washed briefly and transferred into a μ -Slide 18 well Glass Bottom (81817, Ibidi GmbH) with ProLong Glass Antifade Mountant (P36984, ThermoFisher Scientific) before imaging with a confocal microscope (Leica SP8, 20X objective). Primary and secondary antibodies are listed in Table 2.

Multi-photon microscopy analysis

Imaging was conducted by using a multi-photon (MTP) Leica SD 8 microscope configured with 750nm excitation, 3-5% laser power, and a sampling rate of 40 μ s/pixel. Two emission signals with broad spectral ranges were utilized to effectively capture autofluorescent signals. Emission signals were collected in the ranges of 404-506nm and 526-627nm. Furthermore, sampling rates were fine-tuned to enhance image quality. The broad spectral range of the emission signals enabled effective capture of autofluorescent signals, eliminating the need for external dyes. Three timepoints (days 4, 12, and 16) were selected to observe changes in hAO morphology.

Lentivirus production and infection

Human umbilical vein endothelial cells (HUVEC/TERT2) (Evercyte GmgH, Wien - CHT-006-0008) were grown in culture flasks coated with 0.1% gelatin with an endothelial cell growth medium BulletKit (CC-3124, Lonza) supplemented with 10% FBS and 20 µg/ml G418 (ant-gn-5, InvivoGen) at 37°C in a humidified atmosphere of 5% CO₂.

Lentivirus (LV) was generated by transfecting a HEK-293T cell line (ATCC CRL-3216™) with packaging plasmids (pMDLg/pRRE Addgene plasmid #12251; pRSV-Rev Addgene plasmid #12253; pMD2.G Addgene plasmid #12259) and transfer plasmids (pRRLsin.cPPT.PGK-GFP.WPRE Addgene plasmid #12252) complexed with Polyethylenimine Max (49553-93-7, Kyfora Bio Polysciences). Medium was changed 6h after transfection. Supernatants were collected 72h after transfection, filtered through a 0.33µm PVDF filter (SLGV033RS, Millipore), and then ultracentrifuged for 2h at 70,000 g at 4°C. The LV pellet was then resuspended in DPBS at 4°C and stored in aliquots at -80°C. The concentrated LV were titrated on HEK-293T. The percentages of GFP⁺ positive cells were evaluated by flow cytometry 5 days post-transduction, and LV titers were calculated and expressed in transducing unit (TU) ml⁻¹. HUVEC/TERT2 cells were infected with the supernatants at a multiplicity of infection of 5-6, and polybrene (TR-1003, Sigma Aldrich) was added to a final concentration of 8 µg/ml. They were successfully infected with the eGFP-expressing lentiviral vector as demonstrated by confocal microscopy and flow cytometry (Fig. Supplementary 2A). To confirm HUVEC/TERT2 cell line identity as internationally requested, short tandem repeat analysis was performed as previously described (Fig. Supplementary 3A, 3B) [22].

Assembloid generation

Cell ratios and culture conditions were optimized to achieve assembloid formation and growth. Before establishing the highest-yielding culture conditions, the viabilities of single cell populations (i.e., LF, hAO, and GFP⁺-HUVEC/TERT2 cells) in defined final assembloid media were tested (data not shown). In addition, different substrates and culture supports were evaluated. Finally, hAO (p4-6), LF (p4-6) and GFP⁺-HUVEC/TERT2 cells were counted separately, then centrifuged together at 1400 rpm for 5 min, and seeded in 1:1:1 ratio in BIOFLOAT 96-well plates (F202003, faCellitate) without Cultrex UltiMatrix RGF-BME (Fig. Supplementary 2B). The final medium was composed of hAO and HUVEC/TERT2 media in a 1:1 ratio with 1% FBS

(final concentration), and supplemented with 10 µM Rock Inhibitor for the first 48 h. The medium was changed every two days.

Due to the novelty of the culture process, the formation of human airway 3D structures was observed daily and monitored under light microscopy. The first step was the identification of formed compact aggregates facilitating the selection of the best culture techniques. Formed multicellular clusters were subjected to immunofluorescence, immunohistochemistry (H&E and AB-PAS) and scanning electron microscopy (SEM) to identify assembloids. Crucial parameters such as viability, mucus formation, and ciliary beating were investigated at days 8, 16, and 24.

Assembloid viability

Because the viability of a 3D model is crucial to ensure the maintenance of tissue-like functionality and cellular interactions and to thereby enable the accurate modeling of biological processes and responses, we evaluated this important parameter until day 40. Cell viability was assessed using the 3D organoid cell viability assay (AKES081, Assay Genie) following the manufacturers' instructions. This kit relies on the enzymatic reduction of WST-8 tetrazolium salt by mitochondrial dehydrogenases, generating an orange-colored product whose intensity correlates with the number of viable cells, as measured at 450 nm. The pre-assay incubation time ranged from 1 to 4 h. The assay was performed at 8, 16, and 40 days to monitor assembloid viability.

Scanning electron microscopy

SEM was conducted on assembloids that were fixed with 2.5% glutaraldehyde in PBS overnight. Samples were rinsed with PBS and milliQ water for 5 min and then dehydrated in ethanol solutions of 30 - 50 - 70 - 90 -100% in H₂O for five min twice. Hexamethyldisilazane (HMDS) drying was completed by immersing samples in 100% HMDS for 10 min followed by mounting on SEM stubs. On the following day, the samples were sputter-coated (Quorum Tech) and scanned by SEM EVO 10 (Zeiss).

Forskolin-induced swelling assay

To assess the functionality of airway epithelial cells, activation of the cystic fibrosis transmembrane conductance regulator (CFTR) chloride channel was evaluated using the Forskolin-induced swelling (FIS) assay. hAO at different passages and airway assembloids after 16 days from seeding were treated with 20µM Forskolin (#1099, Tocris), freshly diluted in culture medium. hAO were exposed to Forskolin for 72 hours, whereas assembloids were treated for up

to 8 days. Swelling of the organoids and assembloids was monitored over time as an indirect measure of CFTR activity. Images were acquired at regular intervals, and changes in size were quantified using Fiji software (National Institute of Health, US).

Organoid and assembloid size measurement

To determine the size of hAO and assembloids, bright-field images were acquired with a Nikon Eclipse TS100 microscope equipped with a DS-Fi1 Nikon camera. The number, the diameter and the area of organoids and assembloids were measured using Fiji Software (National Institute of Health, US) by two blinded operators. At least three independent experiments were analyzed.

Statistical analysis

All the quantitative data are presented as mean $\pm$ SEM, as indicated in each figure legend with the specific statistical test used. All experiments were repeated at least three times; each performed on samples from different donors. All statistical tests were performed using GraphPad Prism 9 and statistical significance (*p-value*) in figure asterisks mean $*p < 0.05$, $**p < 0.01$, $***p < 0.001$. P values < 0.05 were considered significant, while p values > 0.05 were not reported.

Results

Generation and characterization of human airway organoids

hAO were generated from biopsy specimens obtained from 15 healthy lung donors (Fig. Supplementary 1C). Organoids presented a spherical shape and an inner cavity with a diameter ranging from 30 μm to 450 μm (Fig. 1A). We improved the hAO culture technique by replacing the mechanical dissociation step with an enzymatic protocol that enabled faster growth and larger organoids than described previously (Fig. 1B) [11]. The reliability of enzymatic dissociation was supported by quantitative analysis showing number of organoids, average size, and cell viability, as assessed by MTT assay (Fig. 1C).

Airway signature

The gene expression profiles of hAO derived from distinct donors were similar during respective passages and recapitulated those of distinct airway tissue cell types, thus proving the reproducibility of this culture method. We demonstrated that mRNAs encoding E-cadherin (E-Cad) and E74-like factor 3 (ELF3), biomarkers of generic airway epithelial cells [23–25], were present during passages. Basal cells

were identified by expressions of keratin-5 (KRT5) and tumor protein 63 (p63), which were constant during different passages (Fig. 1D). Club cell marker secretoglobulin family 1A member 1 (SCGB1A1), also known as club cell secretory protein 10 (CC10), was very highly expressed in comparison to other markers, indicating the maintenance of airway integrity and repair of organoids [26]. Furthermore, anterior gradient protein 2 homolog (AGR2), one of the secretory goblet cell markers, was highly expressed. Ciliary markers nephrocystin 1 (NPHP1) and dynein axonemal heavy chain 5 (DNAH5) were present and both were expressed continuously during passages (Fig. 1D). SRY-box 2 (SOX2) and NK2 homeobox 1 (NKX2.1), key markers of lung progenitor cells, were consistently expressed throughout the passages. Interestingly, leucine-rich repeat-containing G-protein coupled receptor 5 (LGR5) was also stably maintained (Fig. 1D). Flow cytometry confirmed airway identity by demonstrating typical surface and intracellular markers such as EpCAM ($92,08 \pm 2,33$), CD271 ($32,86 \pm 5,92$), CD66c ($55,99 \pm 12,73$), tetraspanin-8 (TSPAN8) ($21,07 \pm 12,02$), and forkhead box protein J1 (FOXJ1) ($36,35 \pm 5,42$) that characterize basal, secretory, and ciliated cells (Fig. 1E and Supplementary 1D). The same signature was observed through confocal microscopy, which additionally demonstrated the specific spatial distribution of these cells that closely mimicked *in vivo* epithelial organization (Fig. 2A-C). To further confirm the airway identity of our hAO, an immunohistochemical assay using a monoclonal antibody highly specific to the human lung alveolar type 2 cell constantly yielded negative results (data not shown) [27,28]. Notably, LGR5, an important marker of adult stem cells and critical modulator of their activity [29], co-localized with p63 (Fig. 2D).

Next, we sought to determine whether our hAO were organotypic and thus able to mimic the cellular composition, histoarchitecture, and functionality of the tissue of origin. AB-PAS staining sections of hAO at different passages (p4 to p8) showed acidic and neutral mucins released by secretory cells into the organoid lumen (Fig. 3A). To ensure a spatially-matched evaluation, reduce inter-sample variability, and enhance the reliability of our findings, we used the same histological sections to allow a direct correlation between mucus production and specific cellular markers. As shown in Fig. 3B, basal cells (p63+ and KRT5+) were in the outer epithelial layer of the organoids, while secretory cells (SCGB1A1+) were in the luminal surface and ciliated cells (beta-IV Tubulin+) were in both apical and luminal surfaces.

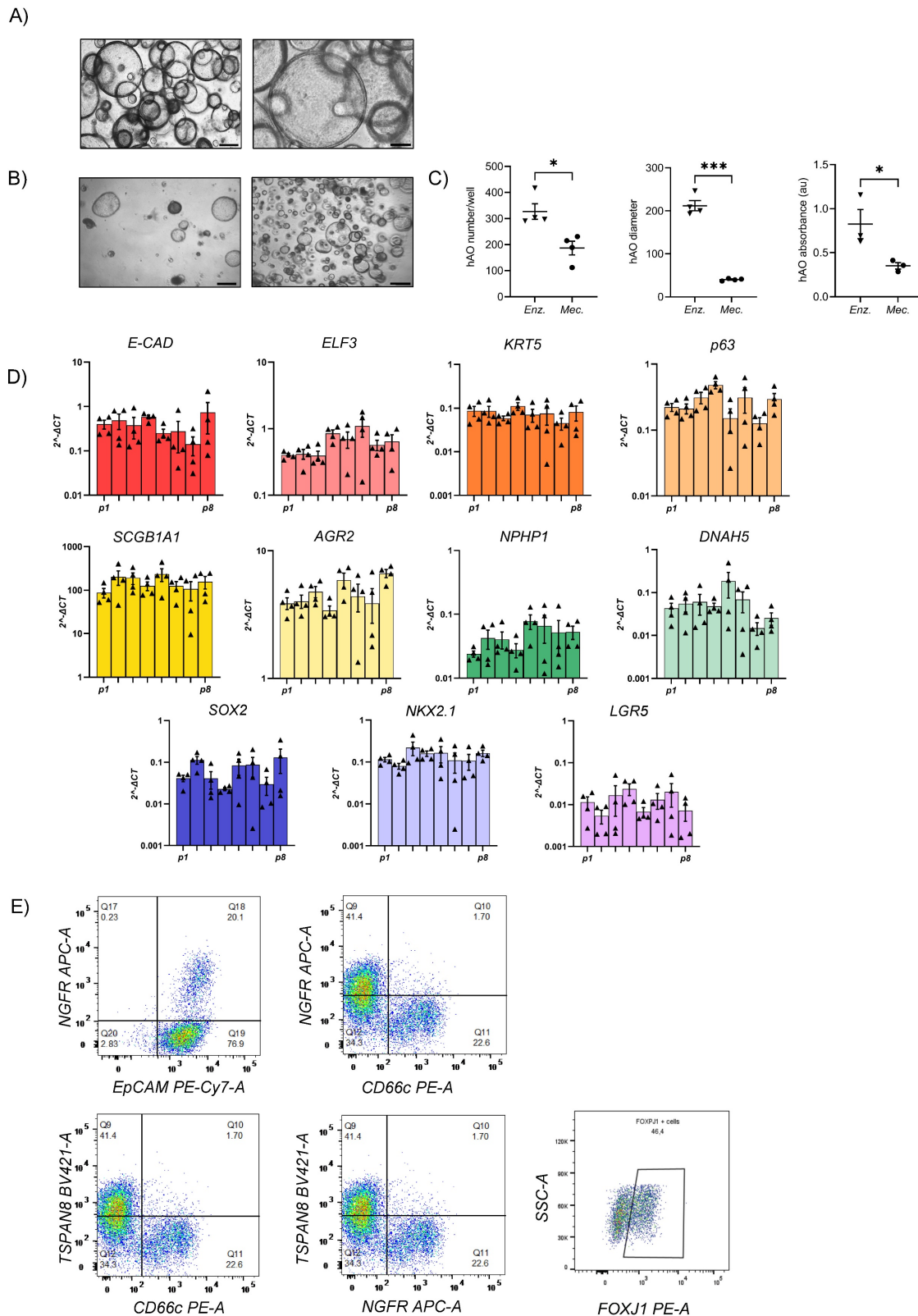


Figure 1. hAO characterization. (A) Representative bright-field images of hAO after 21 days from seeding. Scale bars 200µm (left) and 90µm (right). Images were acquired using a DS-Fi3 Nikon camera; (B) Representative bright-field images of hAO in two different seeding protocols: from mechanically (left) to enzymatic dissociation (right). Images were acquired using a DS-Fi1 Nikon camera. Scale bars 200µm for both images; (C) Quantification of organoid number (left), diameter in µm (middle) and viability (right) comparing the two different seeding protocols. The data reported in the graphs are mean ± SEM (n=4; n=3 for viability). Statistical analysis was performed using two-tailed unpaired t test with Welch's correction for organoid number and diameter analysis (*p<0.05, ***p<0.001); two-tailed unpaired t test for viability analysis (*p<0.05); (D) Gene expression levels of different epithelial markers demonstrated the airway feature of organoids during passages, from p1 to p8. The data reported in the graphs (log₁₀) are mean ± SEM of 2^{-ΔΔCT} of different organoid cell lines (n=4); (E) Representative flow cytometry dot plots of hAO surface and intracellular airway markers at 21 days after seeding (n=6). Antibodies reported in Table 2.

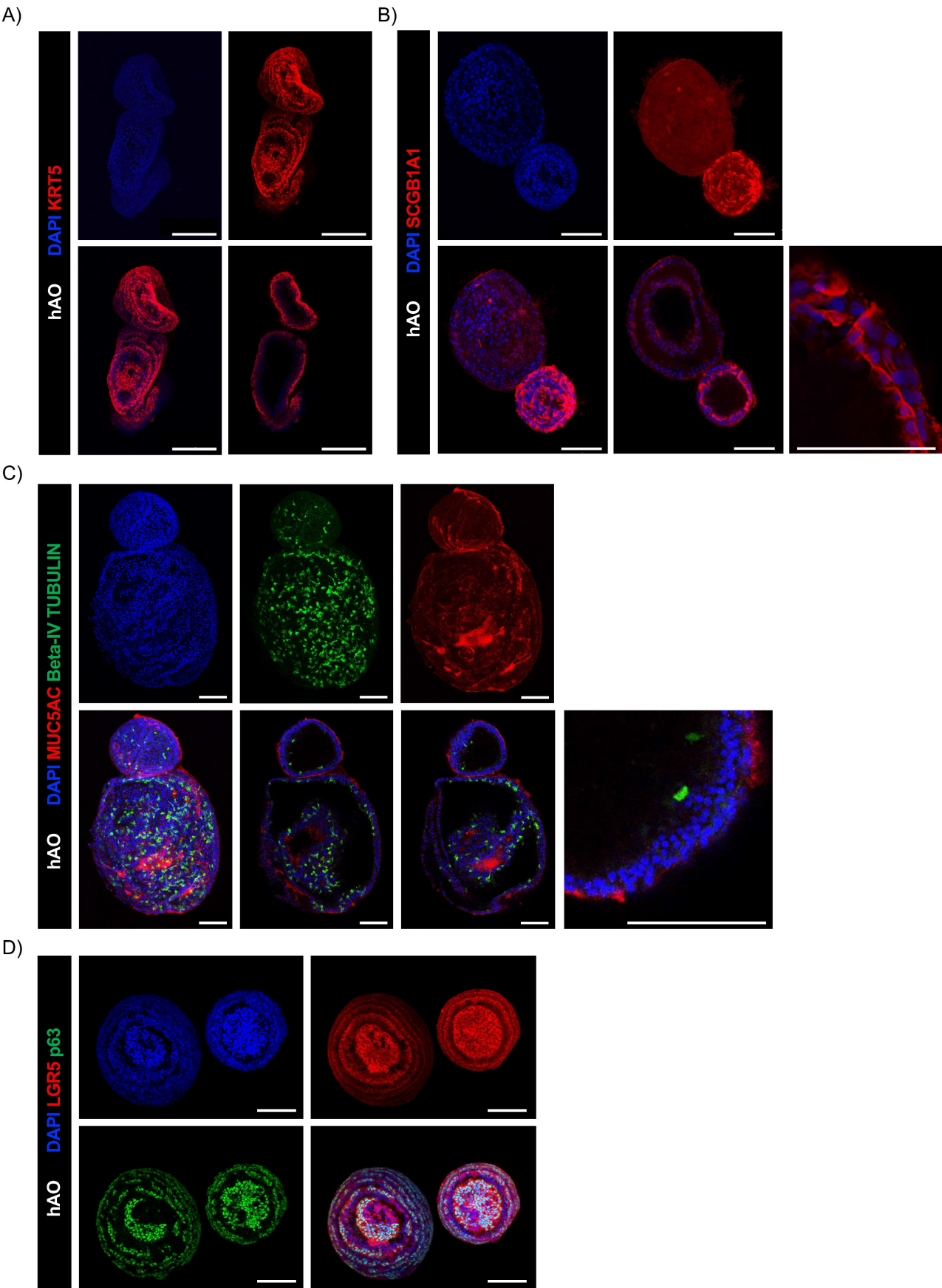


Figure 2. Confocal microscopy images of hAO. Representative confocal microscopy z-stacks and maximum projections of hAO (n=4), objective 20X (Leica SP8). Scale bar 100μm. (A) KRT5 (in Alexa Fluor-555); (B) SCGB1A1 (in Alexa Fluor-555); (C) MUC5AC (in Alexa Fluor-555) and beta-IV Tubulin (in Alexa Fluor-488); (D) LGR5 (in Alexa Fluor-555) and p63 (in Alexa Fluor-488).

MTP microscopy of hAO

MTP microscopy enabled the examination of cellular structures and properties while preserving the organoids' natural environments. To analyze the morphological changes and the modification of cell shapes during growth, we performed an MTP living-analysis. Four days after seeding, hAO displayed a reticular organization, with nuclei encircled by filamentous structures. hAO size ranged between 10-20 μm . On day 12 after seeding, clusters of nuclei formed a circular shape, and hAO sizes ranged from 50 to 100 μm . On day 16, two autofluorescence signals were detected, indicating the presence of specific cell populations. At this stage, the organoids underwent significant morphological changes, growing to sizes of approximately 200-300 μm . Nuclei formed a circular pattern, with a noticeable thickening of the cells located on the organoid exterior. Additionally, the presence of mucin in the central lumen suggested functional differentiation (Fig. 3C, shown in orange).

Generation and characterization of human lung fibroblasts

LF were successfully isolated from the same healthy lung donor biopsy specimens. These primary cell lines were fully characterized by RT-qPCR and flow cytometry that revealed expressions of CD90, CD73, CD105, α -SMA, and vimentin; and negativity for CD45 (Fig. 4A-C and Suppl. 2B). The detection of ECM-specific markers such as COL1A2 and fibronectin by flow cytometry and immunofluorescence staining confirmed the ability of our LF to support the secretion of ECM (Fig. 4B-C).

Generation and characterization of human airway assembloids

To create a model that more closely resembles the natural organ, we developed an assembloid that replicates healthy airway structure. First, we investigated the ability of synthetic ECMs to support the formation of a 3D structure, but failed to establish genuine intercellular communication. Therefore, we used innovative devices to build and support rapid cell aggregation with high reproducibility, and omitted plate centrifugation to reduce cellular stress. We therefore selected a specific low-attachment U-bottom plate coated with a fully xenofree polymer that avidly facilitates cellular interactions. This approach avoided the use of soluble ECM and yielded a scaffold-free system (Fig. Supplementary 2C).

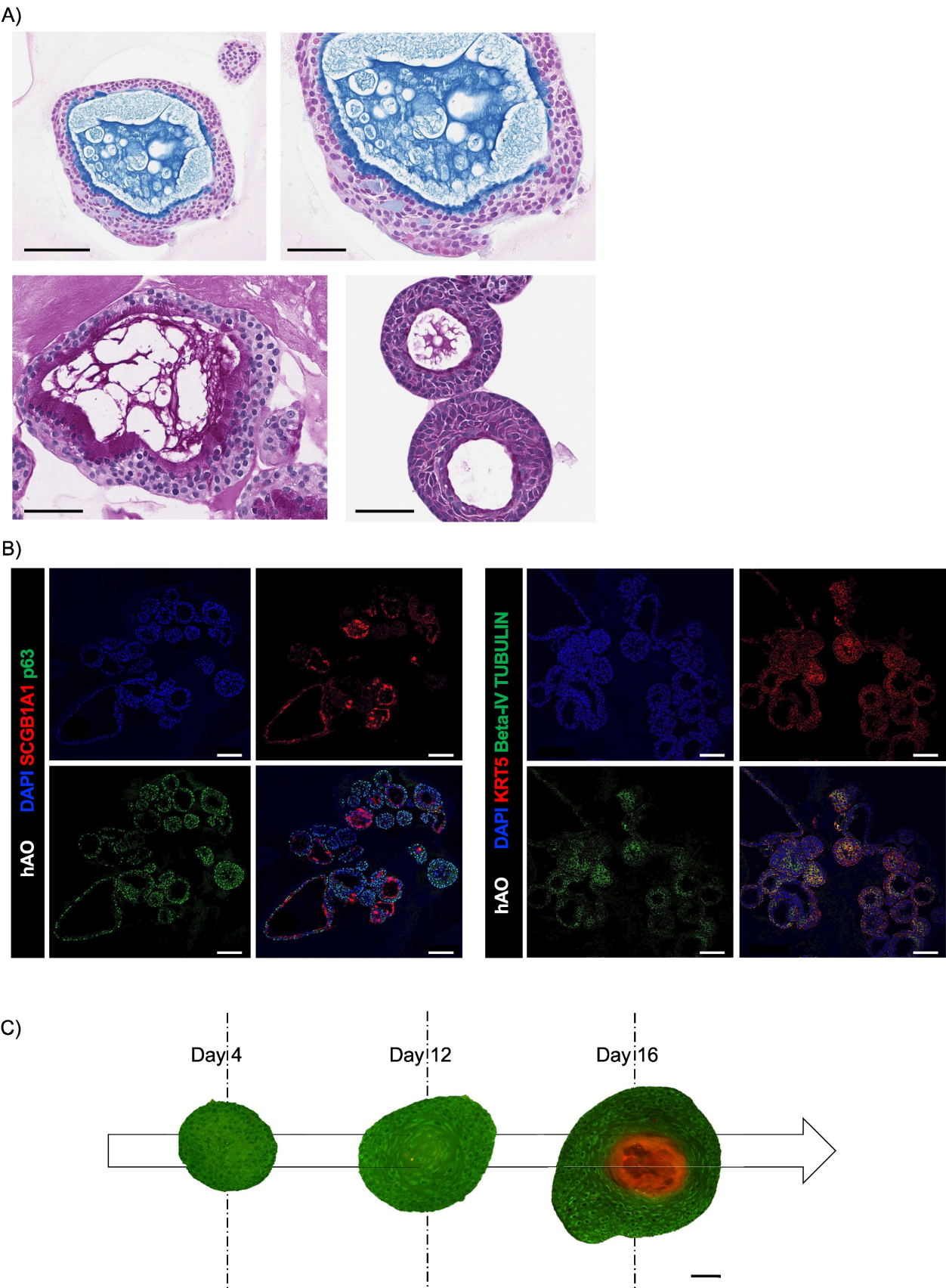
The monitoring of assembloids from initial seeding enabled assessment of the interplay between epithelial and stromal compartments and

investigation of the chronology and dynamics of cluster formation (Fig. 5A). First, we observed that the fibroblast compartment was essential for driving the formation of an organized 3D structure by supporting ECM secretion [30]. In contrast, when only epithelial and endothelial components were present, no structured assembly occurred (Fig. 5B). Confocal microscopy on days 8, 16, and 24 disclosed the resemblance of the assembloid 3D structure to that of natural airways, the interactive organization of differentiated cell types and epithelial function.

Endothelial cells were organized in the luminal surface and were surrounded by stromal cells, while epithelial cells formed an outer layer, resembling a basement membrane (Fig. 5C and Fig. Supplementary 2D). Furthermore, our assembloids remained viable for up to 40 days, maintaining good cell viability, structural integrity, and a well-preserved microenvironment (Fig. 6A).

Summarizing the key functions of the tissue of origin is as relevant as viability to prove that this avatar is not an inactive reproduction but a true mini-organ. The demonstration of mucus production and ciliary movement confirmed assembloid functionality and activity, thus validating that our model is biologically relevant and well-maintained in culture. This was also supported by H&E and AB-PAS staining and by SEM, which, as previously performed for the hAO, exhibited both basal and secretory cells (Fig. 6B-C). Additionally, ciliated cells within cystic airway organoids (Video 1S) and assembloids (Video 2S and Video 3S) disclosed both apical and luminal orientations, facilitating their motility within the ECM and further emphasizing the dynamic nature of the model [31].

To further demonstrate the functionality of the airway epithelial component, we analyzed key epithelial ion channels at both transcriptional and protein levels in hAO. For the ENaC, the α -subunit, indispensable in the maintenance of fluid homeostasis in the lung, was tested [32]. Particularly, CFTR and ENaC- α were stable during passages, from p1 to p8, as demonstrated by gene expression and confocal imaging analysis (Fig. 7A, 7B, Fig. Supplementary 3C). Moreover, to assess the functional activity of CFTR we performed the forskolin-induced swelling assay, which demonstrated a significant increase in organoid size over time, confirming active ion and fluid transport (Fig. 7C). To validate the functional integrity of the airway epithelium in the assembloid, the same forskolin administration was performed and a progressive and significant increase in size over time was observed (Figure 7D).



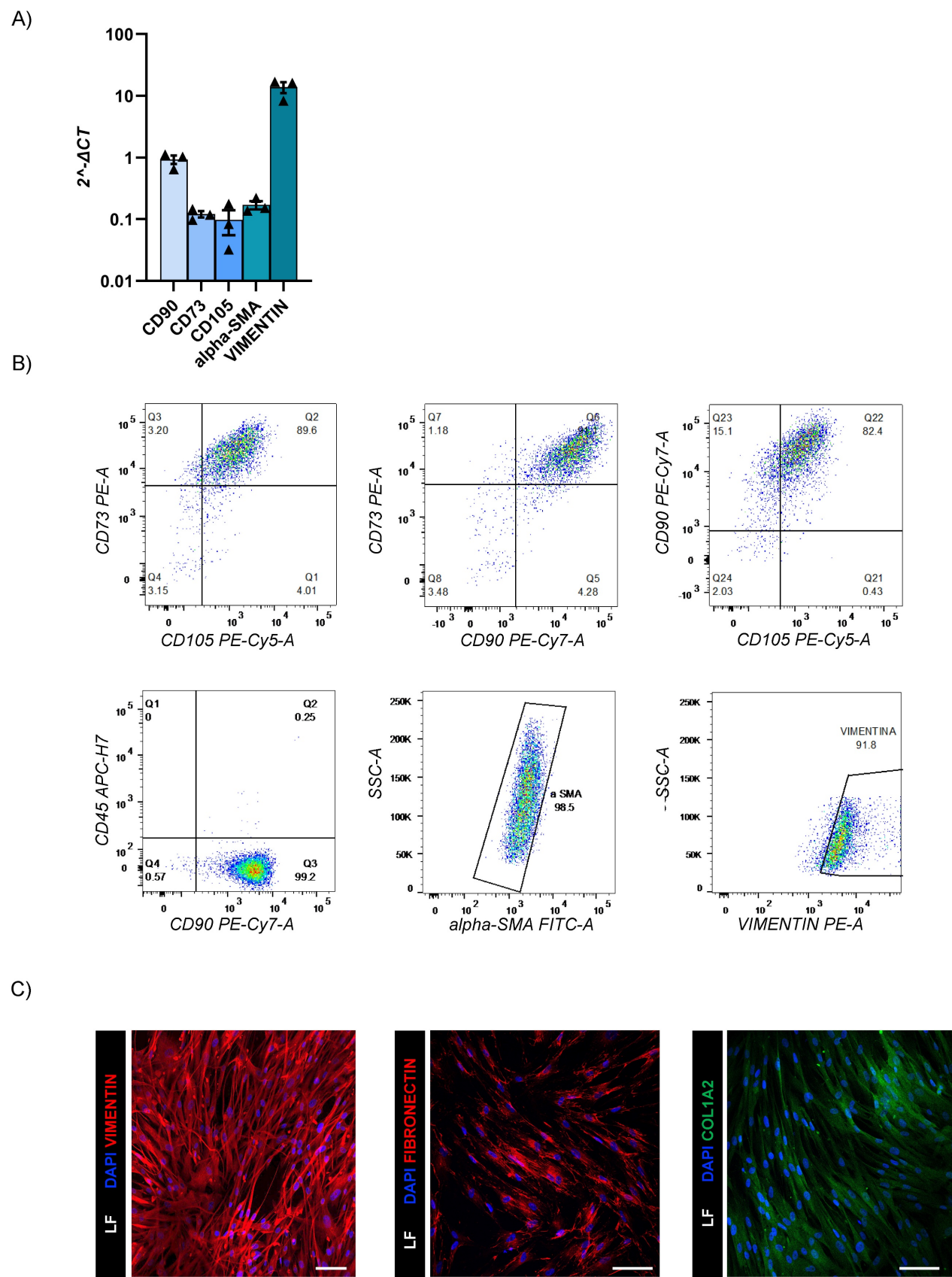


Figure 4. LF characterization. (A) Gene expression levels of fibroblast markers. The data reported in the graphs (log10) are mean $\pm$ SEM of $2^{-\Delta\Delta CT}$ of different LF donors (n=3); (B) Representative flow cytometry dot plots of LF surface and intracellular markers (n=3); (C) Representative confocal microscopy images of intracellular vimentin (Alexa Fluor-555), fibronectin (Alexa Fluor-555) and COL1A2 (Alexa Fluor-488) markers (n=3). Scale bar 100µm (Leica SP8).

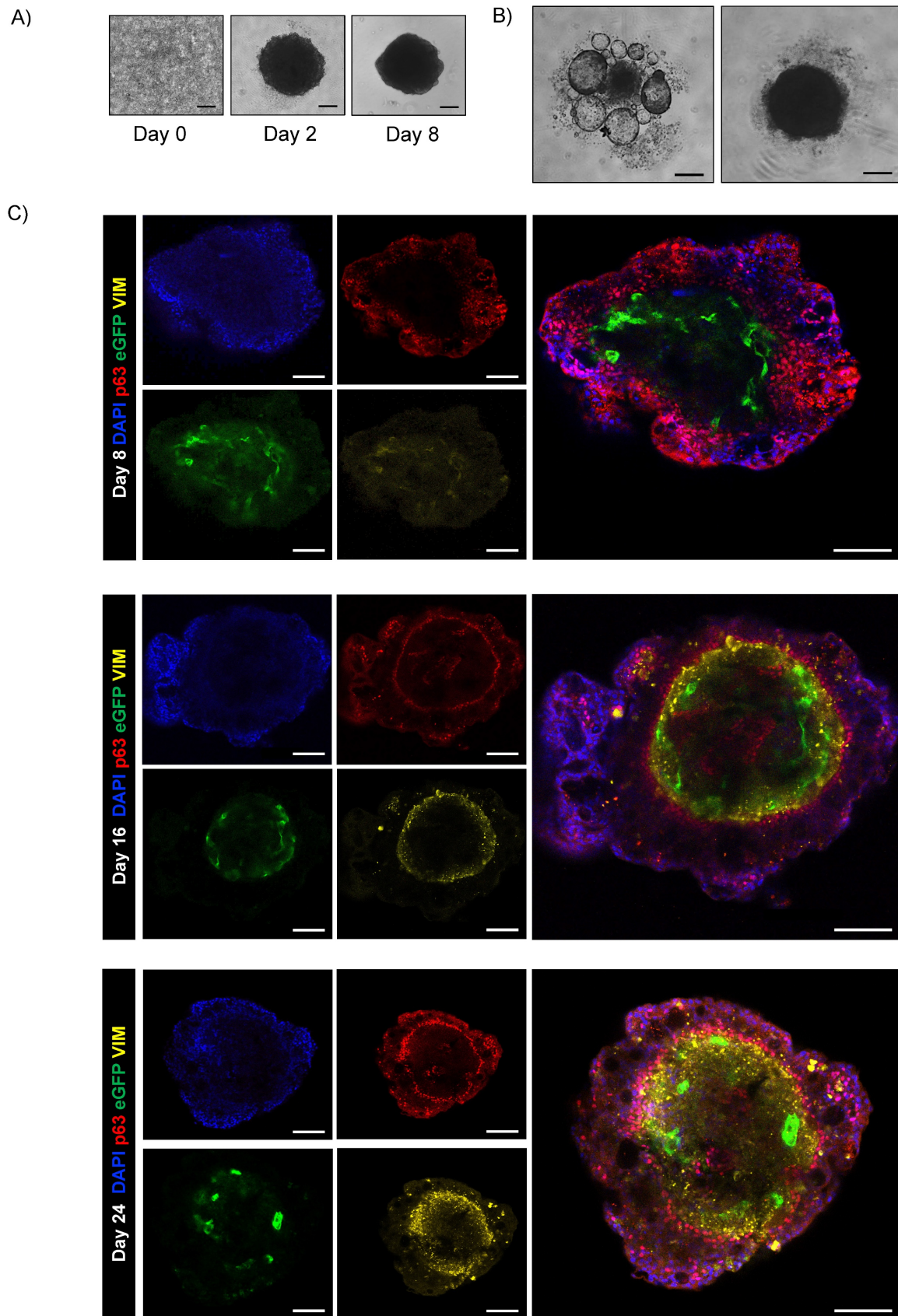
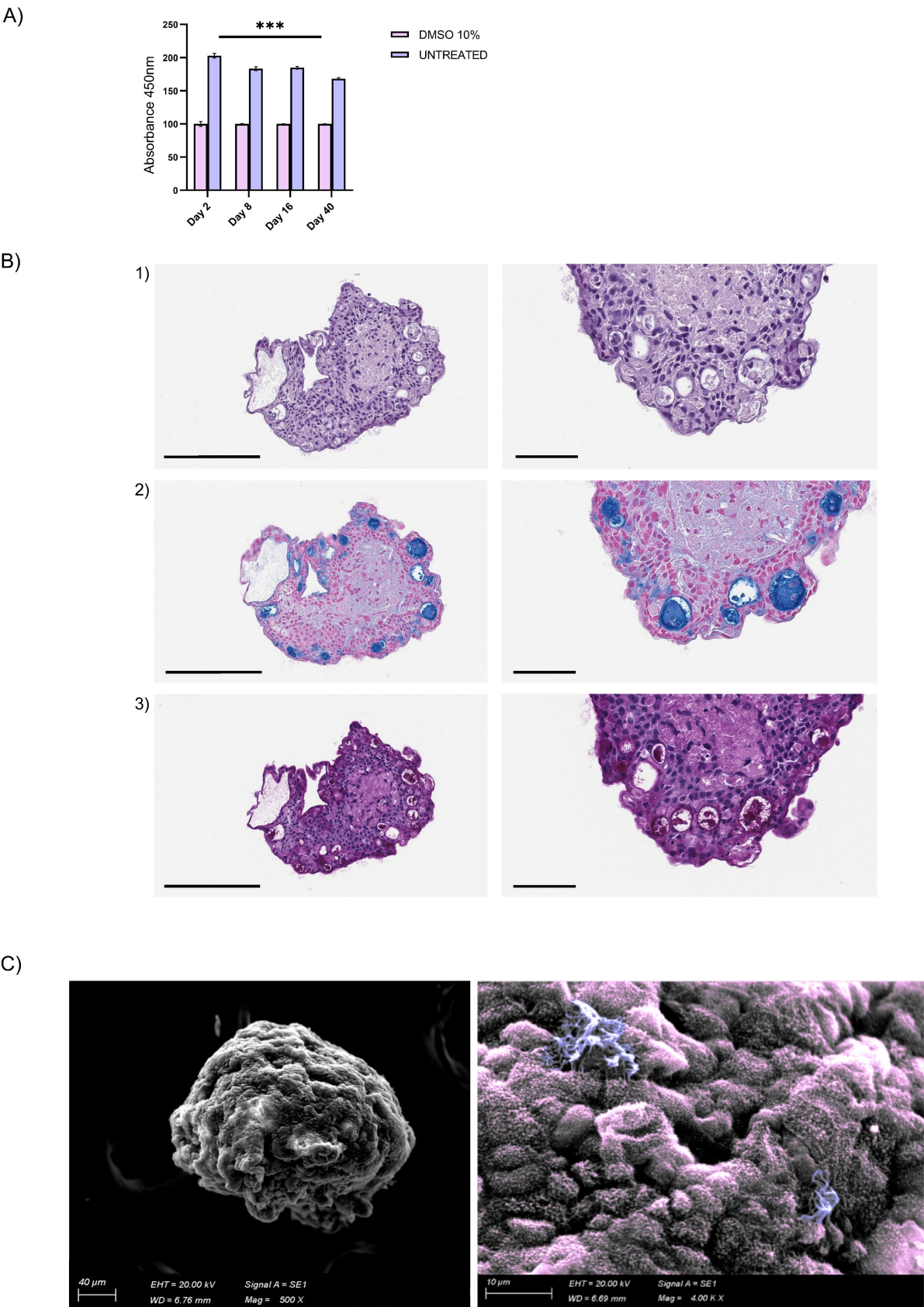


Figure 5. Airway assembloid generation and characterization. (A) Representative bright-field images of human airway assembloids at different days from seeding. Scale bars 200 μ m. Images were acquired using a DS-Fi3 Nikon camera; (B) Representative bright-field images of hAO with HUVEC/TERT2 (left) and hAO with LF (right). Scale bars 200 μ m. Images were acquired using a DS-Fi1 Nikon camera; (C) Representative confocal microscopy images of human airway assembloids at different time points: day 8, day 16 and day 24 (n=6). Basal cells represented by p63 (in red), LF by vimentin (in yellow) and HUVEC/TERT2 eGFP+ (in green). Objective 20X (Leica SP8). Scale bar 100 μ m.



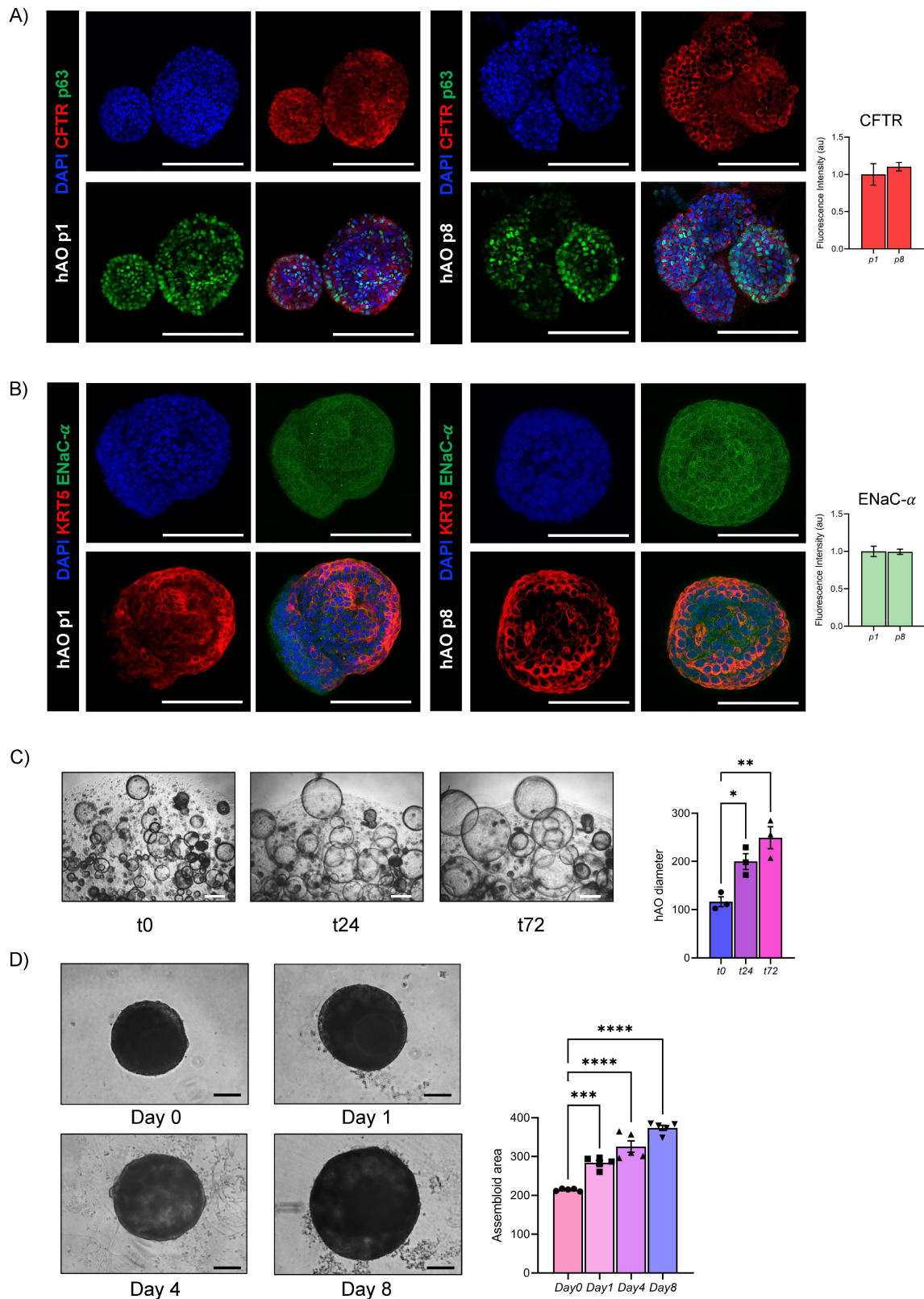


Figure 7. Functional evaluation of airway epithelial physiology. (A) Representative confocal microscopy maximum projections of hAO, objective 20X (Leica SP8). Scale bar 100µm. CFTR (in Alexa Fluor-555) and p63 (in Alexa Fluor-488) (left); quantification of mean fluorescence intensity with background subtraction (right). The data are presented as mean $\pm$ SEM (n=3 at passages 1 and 8). (B) Representative confocal microscopy maximum projections of hAO, objective 20X (Leica SP8). Scale bar 100µm. KRT5 (in Alexa Fluor-555) and ENaC- α (in Alexa Fluor-488) (left); quantification of mean fluorescence intensity with background subtraction (right). The data are presented as mean $\pm$ SEM (n=3 at passages 1 and 8). (C) Representative bright-field images of hAO at t0, t24, t72 after forskolin treatment (20µM) were acquired using a DS-Fi1 Nikon camera. Scale bars 200µm (left). Quantification of organoid diameter (in µm) at different timepoints after treatment (right). The data reported in the graphs are mean $\pm$ SEM (n=3). Statistical analysis was performed using one-way ANOVA, Dunnett's multiple comparison test vs t0 (* p <0.05 t0 vs t24, ** p <0.01 t0 vs t72). (D) Representative bright-field images of airway assembloids at day 0, day 1, day 4, day 8 after forskolin treatment (20µM) were acquired using a DS-Fi1 Nikon camera. Scale bars 200µm (left). Quantification of the assembloid area at different timepoints after treatment (right). The data reported in the graphs are mean $\pm$ SEM (n=3). Statistical analysis was performed using one-way ANOVA, Dunnett's multiple comparison test vs Day 0 (* p <0.05 of t0 vs t24, ** p <0.01 t0 vs t72).

Discussion

New and advanced models will not only elucidate the pathobiology of pulmonary disorders but may also accelerate therapy development. Lung organoids represent easier-to-use and more ethically sustainable alternatives to animal models while closely simulating human pulmonary physiology and preserving the genetic and phenotypic features of their native tissue [33]. Although animal models remain the primary tools for studying the pathobiology of lung diseases such as pulmonary fibrosis, chronic obstructive pulmonary disease, lung cancer, and play a key role in preclinical drug screening, their high cost, lengthy experimental timelines, and variability often lead to poor reproducibility and limited translational potential [34]. Moreover, aligning with the European 3Rs principle (Replacement, Reduction, and Refinement), these models may minimize animal use in research [35–37].

In 2009, Rock et al. used murine adult airway stem cells to establish the first airway organoid culture protocol [38]. A decade later, in 2019, Sachs and colleagues developed a long-term airway organoid culture system [21]. Over the last decade, airway organoid technology has emerged as a versatile research tool for replicating the diversity of human airway pseudostratified epithelia and facilitating the *in vitro* study of neoplastic and infectious pulmonary diseases [39].

In this study, we first optimized the generation and culture of airway organoids derived from donor biopsy specimens, thus advancing our previous work [11]. This methodology, based on enzymatic dissociation, enables the precise counting of single cells that constitute organoids, ensuring reproducibility in the culture system. Our hAO consistently expressed typical markers of basal, secretory (including club and goblet), and ciliated cells during passages, as demonstrated by multiple techniques. Moreover, levels of SOX2 and NKX2.1 indicated the self-renewal capacity of these adult stem cells, promoting their differentiation and supporting the identity and function of epithelial cells.

We detected LGR5 expression, a biomarker of stemness, during hAO passages. LGR5 is essential for self-renewal and differentiation, it is a key marker of active stem cells in organoids and unequivocally identifies self-renewing organoids [13]. Indeed, its expression prolongs the stability of the model and maintains tissue regenerative capacity.

The most crucial indicator of functionality is mucus secretion [40], which was demonstrated not only by standard methods but also through the

application of the most advanced and innovative technologies. Indeed, mucus secretion was further confirmed by MTP microscopy and label-free images that are advantageous because of their preservation of natural cellular structure and function while reducing interference and artifacts from dyes. This ground-breaking approach revealed ultrastructural details that included circular patterns of nuclear organization and sequential mucus secretion, suggesting the functional commitment of the organoids.

Although organoids are the most widely used 3D models in tissue modeling, they still exhibit limitations, particularly in establishing complex interactions between tissue cell types. Limited information regarding vascular structures, their interactions with the microenvironment, and inter-organ signaling continue to compromise their fidelity to human organs [41]. A transition from the use of organoids to more complex networking structures such as assembloids is needed to facilitate studies of the interactions between multiple cell types and to thereby better understand pathobiology.

Assembloids are 3D cell culture systems including complex and integrative models that incorporate multiple types of organoids or specialized cell types. These structures are self-organizing, and thereby create a more comprehensive and dynamic system that better simulates interactions within or between organs. Assembloids are useful for investigating tissue remodeling, complex diseases, and multi-tissue dynamics [42]. Consequently, assembloids are increasingly recognized as a better alternative in multiple fields of biomedical research, and lead the advancement of stem-cell research. Recent studies have focused on assembloid generation through multicellular co-culture and 3D extrusion printing. This technology has been applied to study normal tissues of the brain and gastrointestinal tract [43–46] as well as bladder and small cell lung cancers [47,48].

Based on these advancements, we developed a new 3D model of healthy lung tissue using an experimental methodology that enabled the self-aggregation of hAO in a more complex structure through interactions between primary lung fibroblasts and an endothelial cell line. Organoids typically grow in solid basement membrane extracts, with Matrigel being one of the most used options due to its ECM proteins that include laminin, collagen type IV, proteoglycans, and growth factors. To create a vital, easy-to-use, and simply reproducible model, we generated our avatar without the use of a synthetic ECM. Lung fibroblasts included in these structures secreted ECM proteins such as COL1A2 and

fibronectin. Interestingly, we observed that the stromal component drove appropriate assembloid formation, while the presence of only epithelial and endothelial cells did not generate an organized model. These experiments confirmed the crucial role of stromal cells in the formation of organized models such as assembloids [49,50]. Furthermore, our approach obviated the use of scaffolds, which are disadvantageous in modeling because of their variability, non-human origin, and potential lack of biological relevance in human tissue systems.

The functions of endothelial cells go beyond their role as structural components of vasculature [51]. Our goal was to incorporate endothelial cells into our 3D system to recreate the *in vivo* environment. However, the isolation of endothelial cells from human lung tissue is difficult due to the coexistence of multiple cell types. After several attempts that yielded a high degree of contamination by non-endothelial cells such as fibroblasts as demonstrated previously by Gaskill et al. [52], we decided to use HUVEC, widely recognized for their key role in angiogenesis [53], to promote the vascular signature within the assembloid model. In addition, the used hTERT-transfected HUVEC cell line has been demonstrated to maintain endothelial characteristics at both mRNA and protein levels similar to those of standard primary HUVEC [54].

The functionality of our assembloid is crucial for its validation as a useful airway model. After assembly that incorporated fibroblasts and endothelial cells, our 3D avatar demonstrated expressions of typical airway markers and key functions of the airway epithelium such as ciliary beating observed by microscopy and mucus production demonstrated by AB-PAS staining.

Our selection of the transcription factor p63 to label the epithelial component deserves further comment. This marker was chosen because it identifies basal cells, which are relatively undifferentiated cells of the pulmonary pseudostratified mucociliary epithelium [38,55]. The use of p63 ensured the identification of basal cells and their persistence through multiple passages. This was crucial to our study because basal cells constitute the most important cell type, without which the lung cannot generate other essential cells. In addition, p63-positive cells serve not only as airway stem cells, but also enable assembloid functionality by producing ciliated and mucosecretory cells. The final and certainly crucial parameter of our 3D assembloid is its long-term viability.

To complement the structural characterization of this 3D model and ensure its physiological relevance, we performed a molecular and functional evaluation

of ion transport mechanisms known to be central to airway epithelial integrity and homeostasis.

Particularly, we demonstrated that from early to late passages airway organoids maintain stable gene and protein expression levels and functional activity of CFTR and ENaC- α , thus confirming that ion homeostasis is physiologically preserved in this system [56]. Interestingly, the functional relationship between the ion channels was not altered even when the airway organoids were integrated in the 3D assembloid structure.

In conclusion, we propose a reproducible 3D healthy airway assembloid as an advanced *in vitro* platform for the study of morphology, functions, and cellular interactions within the lung.

Limitations

Although our 3D lung assembloid model represents a significant advance in lung research, some aspects require optimization. The inclusion of immune cells could further enhance its applicability in studying pulmonary inflammation, while more sophisticated vascularization strategies may improve its recapitulation of native lung microcirculation. Additionally, the absence of airflow dynamics and mechanical stimuli, key factors in lung physiology, could be addressed by integrating bioreactor systems. Despite these limitations, our model sustains viability, functionality, and key cellular interactions over time, thus offering a promising tool for studying lung diseases and testing new therapeutic approaches.

Abbreviations

3D: three-dimensional; 2D: two-dimensional; hAO: human airway organoids; RT: room temperature; FBS: fetal bovine serum; LF: lung fibroblasts; SEM: scanning electron microscopy; H&E: hematoxylin and eosin; AB-PAS: Alcian-blue periodic acid Schiff; ECM: extracellular matrix; MTP: Multi-photon; HUVEC: human umbilical vein endothelial cells; LV: lentivirus; GFP: green fluorescence protein; TU: transducing unit; HMDS: hexamethyldisilazane; SEM: standard error of the mean; SCGB1A1: secretoglobin family 1A member 1; CC10: club cell secretory protein; AGR2: anterior gradient protein 2 homolog; NPHP1: ciliary markers nephrocystin 1; DNAH5: dynein axonemal heavy chain 5; SOX2: SRY-box 2; NKX2.1: NK2 homeobox 1; LGR5: leucine-rich repeat-containing G-protein coupled receptor 5; TSPAN8: tetraspanin-8; FOXJ1: forkhead box protein J1; KRT5: keratin-5; p63: tumor protein 63; E-Cad: E-cadherin; ELF3: E74-like factor 3; CFTR: cystic fibrosis transmembrane conductance regulator; α -ENaC: epithelial sodium channel alpha subunit.

Supplementary Material

Supplementary figures.

<https://www.ijbs.com/v21p6234s1.pdf>

Video S1. <https://www.ijbs.com/v21p6234s2.mp4>

Video S2. <https://www.ijbs.com/v21p6234s3.mp4>

Video S3. <https://www.ijbs.com/v21p6234s4.mp4>

Acknowledgments

The authors thank Mr. Marco Brevi, Division of Pathology, Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, Milan, Italy, for his valuable support with immunohistochemical preparations; Dr. Elena Longhi, Laboratory of Transplant Immunology, SC Trapianti Lombardia - NITp, Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, Milan, Italy, for her analysis of Short Tandem Repeats; prof. Loris Rizzello, Department of Pharmaceutical Sciences (DISFARM), University of Milan, Italy and National Institute of Molecular Genetics "Romeo and Enrica Invernizzi" (INGM), Milan, Italy for providing access to the confocal microscopy facility, and finally Dr. Alessandro Olivieri for his precious technical support.

Funding

This work was funded by the European Union - Next Generation EU - NRRP M6C2 - Investment 2.1 Enhancement and strengthening of biomedical research in the NHS (Project title "Exploiting the power of human induced pluripotent stem cell extracellular vesicles as a new anti-inflammatory drug for lung ischemia reperfusion injury", PNRR-POC-2022-12375642 CUP C43C22001430006 I73C22000550006 H73C22001630006). This research was also funded by Ricerca Corrente by the Italian Ministry of Health.

Author contributions

M.C.I. designed and supervised the research. M.C.I. and L.L. wrote, reviewed, and revised the manuscript. M.C.I., A.C., D.T., F.R., F.C., N.E., G.L., M.C., T.D.U., E.S., performed the experiments, and analyzed and interpreted the data. A.D.A., V.M., and L.L. critically revised the manuscript. L.R. provided lung biopsies and revised the manuscript. L.L. provided technical and material support, conceptualized and supervised the experiments. All authors have read, discussed and approved the final manuscript.

Ethical approval and consent to participate

All studies were conducted in accordance with the relevant guidelines and regulations. Written informed consent was obtained from the donors

involved in the study (Figure Supplementary 1C). All experiments involving human cells were carried out following the principles of the Declaration of Helsinki. Ethical evaluation was performed by the Ethical Committee of Milan Area 2 n°6851, February 14, 2023.

Competing Interests

The authors have declared that no competing interest exists.

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