Curcumin/poly(2-methyl-2-oxazoline-b-tetrahydrofuran-b-2-methyl-2-oxazoline) formulation: An improved penetration and biological effect of curcumin in F508del-CFTR cell lines

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Curcumin/poly(2-methyl-2-oxazoline-b-tetrahydrofuran-b-2-methyl-2-oxazoline)

formulation: An improved penetration and biological effect of curcumin in F508del-

CFTR cell lines

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Running title: Curcumin/triblock copolymer and F508del-CFTR cells

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Abstract

Neutral amphiphilic triblock ABA copolymers are of great interest to solubilize hydrophobic drugs. We reported that a triblock ABA copolymer consisting of methyl-2-oxazoline (MeOx) and tetrahydrofuran (THF) ($\text{MeOx}_6\text{-THF}_{19}\text{-MeOx}_6$) (TBCP2) can solubilize curcumin (Cur) a very hydrophobic molecule exhibiting multiple therapeutic effects but whose insolubility and low stability in water is a major drawback for clinical applications. Here, we provide evidences by flow cytometry and confocal microscopy that Cur penetration in normal and ΔF508 -CFTR human airway epithelial cell lines is facilitated by TBCP2. When used on ΔF508 -CFTR cell lines, the Cur/TBCP2 formulation promotes the restoration of the expression of the CFTR protein in the plasma membrane. Furthermore, patch-clamp and MQAE fluorescence experiments show that this effect is associated with a correction of a Cl^- selective current at the membrane surface of F508del-CFTR cells. The results show the great potential of the neutral amphiphilic triblock copolymer $\text{MeOx}_6\text{-THF}_{19}\text{-MeOx}_6$ as carrier for curcumin in a Cystic Fibrosis context. We anticipate that other $\text{MeOx}_n\text{-THF}_m\text{-MeOx}_n$ copolymers could have similar behaviours for other highly insoluble therapeutic drugs or cosmetic active ingredients.

1. Introduction

Many chemotherapeutic drugs or cosmetic active ingredients are very insoluble in water compromising their effectiveness and clinical applications. Delivery of drugs into the cells relies on many constraints such as solubilization, endocytosis or crossing through the membrane lipid bilayer. In this context, more particularly, polymer-based synthetic vectors offer advantages such as relative simplicity of production, safety and versatility. Polyester/ether ABA triblock copolymers developed as drug delivery system comprise poly(D,L-lactide-block-ethylene oxide-block-D,L-lactide) (PLA-PEO-PLA), poly[(D,L-lactide-co-glycolide)-block-ethylene oxide-block-(D,L-lactide-co-glycolide)] (PLGA-PEO-PLGA) and poly(ϵ -caprolactone-block-ethylene oxide-block- ϵ -caprolactone (PCL-PEO-PCL) [1]. PPO-PEO-PPO copolymers, Pluronic polymers, have been studied quite extensively thanks to the formation of core-shell micelles comprising the polyethylene oxide (PEO) block as the hydrophilic shell of the corona and the polypropylene oxide (PPO) block as the core [2, 3]. Such copolymers have already received large attentions to solubilise hydrophobic molecules for drug delivery [4, 5]. Certain Pluronic polymers have shown capacity to pass the blood-brain barrier [2]. They have even demonstrated their ability to increase the biodistribution after oral delivery of molecules poorly soluble in water such as genestein [6]. In a clinical phase I study, doxorubicin bound to a pluronic polymer has demonstrated an increased efficacy compared to free doxorubicin [7]. In the field of immunization with proteins and peptides, neutral amphiphilic copolymers have increased significantly the humoral and cellular responses after intravenous injection [8]. Lutrol has provided efficient gene expression in lung and skeletal muscles upon intratracheal and intramuscular co-injection in mice of plasmid DNA, respectively [9, 10]. Moreover, DNA vaccination has been obtained with DNA/amphiphilic block copolymer nanospheres [11].

Poly(2-oxazolines) based polymers are also interesting biomaterials to solubilize water insoluble molecules [12]. Formulations based on poly(2-oxazoline) polymeric micelles and the effect of parameters related to the structure, formulation, additives and toxicity have been described [4, 13-16]. These reported properties allowed to conclude to the high potential of poly(2-oxazoline) block copolymers, for instance in cancer treatments. Poly(2-methyl-2-oxazoline-b-2-butyl-2-oxazoline-b-2-methyl-2-oxazoline) (p(MeOx-b-BuOx-b-MeOx)) has been found to solubilize high quantity of paclitaxel, a very low soluble molecule in aqueous media exhibiting powerful antineoplastic agent, and demonstrated an improved therapeutic effect in xenograft mice tumor models [17, 18]. A clinical phase I study with a rotigotine polyoxazoline conjugate has been initiated from data obtained in Parkinson's disease [19].

Recently, we have reported the synthesis and characterization of an original family of poly(2-methyl-2-oxazoline-b-tetrahydrofuran-b-2-methyl-2-oxazoline) (MeOx-THF-MeOx) neutral amphiphilic triblock copolymers and showed that some of them solubilized curcumin [20]. Curcumin (diferuloylmethane) (Cur), the natural component of the plant "*Curcuma Longa*" exhibits multiples therapeutic effects but is very insoluble in water [21]. This is a safe drug even at high doses, but is rapidly metabolized and poorly absorbed by the cells [22]. Curcumin is proposed for treatment of various pulmonary diseases including Asthmas, cancer and Cystic Fibrosis [23]. In order to evidence the interest of curcumin solubilisation by MeOx-THF-MeOx copolymers, we decided to investigate its influence in a Cystic Fibrosis context. Indeed, curcumin has been proposed to treat Cystic fibrosis (CF), the most lethal genetic disease caused by mutations in the gene encoding the Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) Cl⁻ channel resulting in abnormal chloride transport at the plasma membrane of many tissues and organs [24]. ΔF508 is the most common mutation of CFTR causing a misfolding of the CFTR protein and its retention in the endoplasmic reticulum (ER) for subsequent proteolytic degradation by the ubiquitin-proteasome pathway

[25]. Cur can partially compensate the ER retention of the defective CFTR protein, both *in vitro* in appropriate cell lines and *in vivo* in F508del-CFTR mice [26, 27]. However, the insolubility and poor stability of Cur in water explain some disappointing results observed in mice treated with Cur [28]. Thus, it appeared for us that curcumin was a good example to test the potential of our amphiphilic copolymers to carry hydrophobic molecules.

Here, we report that the MeOx₆-THF₁₉-MeOx₆ (TBCP2) copolymer facilitates penetration of curcumin in F508del-CFTR cells and preserves the capacity of curcumin to restore a functional CFTR protein.

2. Material and Methods

All reagents were purchased from Sigma (St. Quentin Fallavier, France) unless otherwise stated.

2.1. Polymer synthesis.

Poly(2-methyl-2-oxazoline) – *b* – poly(tetrahydrofuran) – *b* – poly(2-methyl-2-oxazoline) triblock copolymer (TBCP2). The copolymer was synthesized as described [20]. Briefly, to a 250 mL reaction flask containing 236 mmol (17 g) of dry THF, 13.9 mmol (3.91 g) of trifluoromethanesulfonic anhydride (Tf₂O) was added at -9°C. The reaction mixture was stirred during 15 minutes and the polymerization was quenched by adding 55.6 mmol (4.7 g) of MeOx at -9°C. Evaporation of residual THF under reduced pressure yielded to a pTHF prepolymer as a white solid. This solid was dissolved in 40 mL of dry acetonitrile and the temperature was increased to 80°C. MeOx (88.2 mmol; 7.5 g) was added and the solution stirred for 24 hours. The reaction was quenched by 4 mL of 2 M sodium carbonate solution and stirred for 1 hour at room temperature. The copolymer was dried and the crude yield was 80%. A chloroform/water extraction was then conducted, the organic phase evaporated and dried for 2 days under vacuum. The triblock copolymer had hydroxyl and ester end functions (47%) as determined by ¹H NMR. Molecular weight of TBPC2 determined by ¹H NMR was

2400 g.mol⁻¹ with a p(THF) block containing 19 monomers and 2 p(MeOx) blocks of 6 monomers each.

Rhodamine-labelled TBCP. First the amine terminated copolymer (diamino-TBCP) was synthesized. The α,ω -diamino-poly(2-methyl-2-oxazoline) – block - poly(tetrahydrofuran) - block - poly(2-methyl-2-oxazoline), was prepared as above but the polymerization was quenched by addition of a large amount of 1,3-diaminopropane instead of MeOx. Molecular weight of diamino-TBCP was 2200 g.mol⁻¹ with a central p(THF) block of 12 monomers and 2 external p(MeOx) blocks containing 8 monomers each. Despite these variations we have considered that TBCP2 and diamino-TBCP have similar physicochemical behaviours. Rhodamine-labelled TBCP was prepared by reaction of 33 mg diamino-TBCP in 1.5 ml of 0.1 M carbonate bicarbonate buffer, pH 9.3 with 14 mg (~2.6 equivalents) of N-hydroxysuccinimide ester activated rhodamine (Invitrogen) dissolved in 0.5 ml dimethyl sulfoxide. The mixture was stirred at room temperature during 24 hours in the dark. A silica gel thin layer chromatography in chloroform/methanol (1/1 vol : vol) revealed the presence of dye bound to the polymer that did not migrate; the fluorescent polymer did not react with ninhydrin. Then, the polymer was purified by precipitation in dichloromethane; the coloured precipitate was washed in ethyl ether and dried under reduced pressure.

2.2. Curcumin solubilization

Curcumin was purified by crystallisation in hot ethanol. Curcumin (10 g) was solubilized in 20 mL pure ethanol under reflux. The hot solution was filtrated, the filtrate was left to cool down and the Cur was collected by filtration (yield 60 %). A 0.2% TBCP2 solution was prepared by adding 5 ml H₂O to 10 mg TBCP2 and vigorous agitation with vortex until complete dissolution. Then 2 mg of Cur were added to the TBCP2 solution, the Cur/TBCP2 (400:2000; $\mu\text{g}:\mu\text{g}$) mixture was vortexed and then sonicated for 5 min at 20°C at

37 kHz. The solution was then clarified by centrifugation (14,100 x g for 10 min) to remove any precipitate.

2.3. Cells and cell culture

The CFBE41o- human bronchial epithelial (homozygous for the F508del mutation) [29] and ΣCFTE29o- human tracheal epithelial (homozygous for the F508del mutation) [30] cell lines carrying the F508del *CFTR* mutation (so called CF cells) and the 16HBE14o- normal human bronchial epithelial cell line [31] (generous gifts from Dieter Gruenert, San Francisco, CA, U.S.A.) were cultured at 37°C in a 5% CO₂-humidified incubator in 20 ml MEM with Earle's Salts (PAA Laboratories), 0.4% Penicillin (40 Units/ml)/Streptomycin (40 µg/ml) (PAA Laboratories), 2mM L-Glutamine (PAA Laboratories), supplemented with 10% heat-inactivated fetal bovine serum (PAA Laboratories). Tissue culture plastic wares (75 cm²) were coated for 20-30 min at 37°C with MEM with Earle's Salts containing fibronectin (0.01 mg/ml), collagen (0.03 mg/ml) and bovine serum albumin (BSA) (0.1 mg/ml). The culture medium was changed every 2 days. The absence of mycoplasma in cell cultures was determined by using MycoAlert® Mycoplasma Detection Kit (Lonza, Levallois Perret, France).

2.4. Curcumin uptake

Two days prior to the experiments, 1.4×10^4 cells were seeded in 2 cm² well of a 4-well plastic culture plate. For curcumin uptake, cells were incubated at 37°C with the curcumin/ TBCEP2 solution (stock solution: 1.1 mM Cur in 0.2% TBCEP2) at various concentrations. Then, the cells were washed twice with PBS, harvested by trypsin, centrifuged (800 x g for 5 min at 4°C) and suspended in cold PBS. The cell-associated fluorescence intensity was measured with a flow cytometer (FACSort, Becton Dickinson; λ_{ex} = 488 nm; λ_{em} = 530/30 nm) before and after treatment with trypan blue (a final concentration of 0.004

%) in order to quench the extracellular fluorescence of curcumin. The fluorescence intensity was expressed as the mean value of the fluorescence intensity (MFI) of 10,000 cells.

2.5. Immunofluorescence assays

Immunofluorescence was carried out with the following antibodies: mouse anti-human CFTR antibodies (Clone M3A7, LifeSpan Biosciences) (dilution 1/50), rabbit anti-calreticulin antibodies (LifeSpan Biosciences) (dilution 1/100), goat anti-calnexin antibodies (C20, Santa Cruz Biotechnology) (dilution 1/100), goat anti-ERGIC-53 antibodies (A-18, Santa Cruz Biotechnology) (dilution 1/100), Alexa Fluor 568 goat anti-mouse antibodies (Invitrogen) (dilution 1/200), Alexa Fluor 568 donkey anti-goat antibodies (Invitrogen) (dilution 1/200), Cy5 sheep anti-mouse antibodies (Jackson ImmunoResearch) (dilution 1/100).

2.6. Confocal microscopy

The cells (1.4×10^4) were seeded on glass coverslips (14 mm diameter) in 2 cm² well of a 4-well plastic culture plate. Intracellular locations were performed by immunofluorescence. The cells were washed several times with ice-cold PBS, fixed in cold methanol (90% in PBS) for 30 min at -20°C, washed several times with PBS containing 0.5% bovine serum albumin (PBS-0.5%BSA). Coverslips were incubated with primary antibodies in PBS-1%BSA for 1 h at 20°C, washed with PBS-0.5%BSA and then incubated for 45 min at 20°C with secondary fluorescent antibodies in PBS-1%BSA. After several washes with PBS, coverslips were mounted in Vectashield (Vector Laboratories, Inc., Burlingame, CA, USA). Confocal laser scanning microscopy (CLSM) was performed using a Zeiss Axiovert 200M microscope coupled with a Zeiss LSM 510 scanning device (Carl Zeiss Co. Ltd., Jena, Germany). The inverted microscope was equipped with a Plan-Apochromat 63X objective (NA=1.4). The fluorescence was measured at either 560 nm upon excitation at 543 nm (Alexa Fluor 568), or 660 nm upon excitation at 633 nm (Cy5).

2.7. Electrophysiological measurements

The cells were cultured on a glass coverslip that was transferred to the experimental chamber of an upright microscope (BX51WI, Olympus Corporation, Tokyo, Japan). Patch-clamp experiments were performed at 21-23°C. Cells were placed in continuously flowing (1-2 ml/min) bath solution containing (mM): 150 NaCl; 6 CsCl; 1 CaCl₂; 1 MgCl₂; 10 D-glucose; 10 HEPES (adjusted to pH 7.4 with Tris) and identified at 60x magnification with a CCD camera (XC-ST70CE, Sony). Somatic whole-cell recordings were performed as previously described [32]. Briefly, low resistance (4-6 MΩ) patch-pipettes pulled on a vertical puller (PB-7, Narishige, Tokyo, Japan) from borosilicate capillaries (Clark Electromedical Instruments, Edenbridge, UK) were filled with internal solution containing (mM): 100 L-aspartic acid; 94 CsOH; 26 CsCl; 14 NaCl; 1 MgCl₂; 3 MgATP; 1 EGTA; 10 HEPES (adjusted to pH 7.3 with Tris). Signals were amplified using the MultiClamp 700B amplifier (Axon Instruments, Foster City, CA). Series resistance was monitored continuously and was typically compensated by 60-70 % in whole-cell configuration. Voltage-clamp recordings were filtered at 4 kHz, sampled at 10 kHz using a data acquisition board (Digidata 1322A, Axon Instruments) operated by Pclamp10 software (Axon Instruments). Off-line analysis was performed using Clampfit10 (Axon Instruments) and Origin8 (Origin Lab Corporation, Northampton, MA). 16HBE14o- and ΣCFTE29o- cells were cultured under control conditions or in the presence of Cur/TBCP2 treatment. Currents were elicited by voltage steps applied from -110 mV to +70 mV (10 mV increments, 400 ms duration) from a holding potential (HP) of -40 mV. Steady-state current amplitude was measured at the end of the pulse and normalized to the cell membrane capacitance (17.6 ± 1.5 pF, n=19 on 16HBE14o- cells; 22.1 ± 1.2 pF, n=28 on ΣCFTE29o- cells).

2.8. MQAE fluorescence assay

The Cl⁻ channel activity of CFTR was assessed on 16HBE14o- and CFBE41o- cells using the halide-sensitive fluorescent probe MQAE [33]. Cells were loaded with MQAE

intracellular dye by incubation for 8 min at 37°C in a hypotonic medium (mM) (110 NaI, 1.92 K₂HPO₄, 0.64 KH₂PO₄, 8 HEPES, 0.8 CaSO₄, 8 D-Glucose, pH 7.4) containing 10 mM MQAE. Coverslips were mounted on the stage of an inverted microscope (LEICA DMI 6000B) equipped for fluorescence, and cell fluorescence was excited at 380 nm. The emitted fluorescence was detected at 470 nm by a CCD camera coolsnap HQ2 (Roper). Cells were maintained at 37°C and continuously perfused with an extracellular bath solution containing (mM): 138 NaI, 2.4 K₂HPO₄, 0.8 KH₂PO₄, 10 HEPES, 1 CaSO₄, 10 D-Glucose, pH 7.4. A superfusion system (Biosciences Tools) allowed rapid change of different extracellular experimental media. Cells were sequentially perfused with 138 mM I⁻ buffer solution, 138 mM NO₃⁻ buffer solution, 138 mM NO₃⁻ buffer solution with 0.5 mM 8-(4-chlorophenyl)thio-cyclic AMP (8-CPT-cAMP) and again with 138 mM I⁻ buffer solutions. Single cell fluorescence intensity was plotted against time at an acquisition rate of 1 image *per* 15 seconds. Fluorescence intensity was normalized to the initial fluorescence level measured in the presence of I⁻.

2. Results

3.1. Curcumin solubilisation and penetration into CF and normal airway epithelial cell lines

The MeOx₆-THF₁₉-MeOx₆ ABA copolymer (TBCP2) of 2400 g.mol⁻¹ was composed of 12 poly(2-methyl-2-oxazoline) (MeOx) hydrophilic blocks A and 19 polytetrahydrofuran (THF) hydrophobic blocks B (Fig. 1A). The chemical shifts of the pTHF blocks were at 1.61 ppm and 3.40 ppm, those of the pMeOx ones at 2.13 ppm (CH₃ groups) and 3.42 ppm and those of the telechelic CH₂ groups adjacent to the end hydroxyl functions at 3.78 ppm (Fig. 1C). The surprising high dispersity ($\bar{D}$ = 4.5) was attributed to the low molar mass of the pTHF block associated to the reversible cationic ring opening polymerization of THF. The standard conditions for MeOx polymerization allowed expecting a much better control of these blocks synthesis as previously reported [34]. The amphiphilic feature in aqueous

241 solution and self-assembling to form micelles of TBCP2 were determined by different
242 techniques including Nile Red fluorescence spectroscopy (F), isothermal titration calorimetry
243 (ITC), dynamic light scattering (DLS) [20]. The critical micellar concentration was 3.2×10^{-3}
244 mol.L^{-1} , $6.5 \times 10^{-4} \text{ mol.L}^{-1}$, $4.2 \times 10^{-4} \text{ mol.L}^{-1}$ to $1 \times 10^{-3} \text{ mol.L}^{-1}$ as determined by F, ITC and
245 DLS, respectively. In a 0.2% solution in water, TBCP2 formed micelles with a hydrodynamic
246 diameter of $16 \pm 1.4 \text{ nm}$ as determined by DLS (Fig. 1D). The solubilisation of 2 mg Cur (400
247 $\mu\text{g/ml}$; 1.1 mM) was achieved by mixing, vortexing and then sonicating 2 mg Cur in 5 ml of
248 0.2% TBCP2 in water (Fig. 1B). In contrast, Cur was completely insoluble in water in the
249 absence of TBCP2 (Fig. 1B). The resulting Cur/TBCP2 formulation (17 wt.%) formed
250 nanoparticles of $255 \pm 30 \text{ nm}$ (Fig. 1D).

251 We then tested whether the Cur/TBCP2 formulation allowed for Cur uptake by the
252 cells. For this purpose, two CF ($\Sigma\text{CFTE29o-}$ and CFBE41o-) and one normal (16HBE14o-)
253 human airway epithelial cell lines were incubated in complete culture medium at 37°C in the
254 presence of Cur solubilized in TBCP2 or TBCP2 without Cur. After 2h incubation, Cur
255 location was analysed by fluorescence microscopy thanks to Cur fluorescence at 520 nm upon
256 excitation at 488 nm. As shown in Figure 2, the fluorescence was in the cytoplasm with
257 Cur/TBCP2 showing penetration of Cur in these cell lines. In contrast, no fluorescence was
258 detected in the cells incubated with TBCP2 in the absence of Cur or the supernatant of Cur in
259 PBS (Fig. 2). The Cur fluorescence intensity associated with the cells was measured by flow
260 cytometry after 2h incubation with various dilution of the Cur/TBCP2 formulation. As shown
261 in Figure 3, the mean fluorescence intensity (MFI) increased with the Cur concentration.
262 Those MFI corresponded to the fluorescence of Cur. Indeed, MFI of the three cell lines
263 incubated with TBCP2 at the concentration used to solubilize $220 \mu\text{M}$ Cur were the same as
264 MFI (2.5 A.U) measured for those cells incubated in the medium without any polymer and
265 Cur. MFI were reduced after treatment with trypan blue which quenched the extracellular

curcumin fluorescence and thus the residual MFI were indicative of Cur internalization. The amounts of Cur associated with the three cell lines and that taken up by the three cell lines were comparable. About ~55% of Cur associated with the cells was inside the cells. For information, MFI upon trypan blue treatment of Σ CFTE29o-, 16HBE14o- and CFBE41o-cells incubated with 27.5 μ M Cur solubilized in DMSO were 45, 13 and 18, respectively, indicating that the penetration of Cur was larger than Cur solubilized in TBCP2 micelles. We evaluated by confocal microscopy the uptake of the copolymer by incubating those cell lines for 4h at 37°C with a Cur/Rho-TBCP formulation (Fig. 4). The images showed that the copolymer did not enter deeply into the cytoplasm of Σ CFTE29o- cells while curcumin enters the cells. It was mostly remained close to the cell surface either at and/or under the cell surface while a few amount of polymer was inside the cells. Regarding the bronchial epithelial cells (CFBE41o-and 16HBE14o- cells), the presence of yellow spots in the cytoplasm was indicative of higher TBCP internalization than in the tracheal epithelial cells (Σ CFTE29o- cells). Next, the evaluation of the effect of curcumin on CFTR in F508del-CFTR cells was studied. The CFTR intracellular location was performed by immunofluorescence and confocal microscopy analysis, and its functionality (assessed *via* chloride current measurements) by patch-clamp experiments and MQAE fluorescence assay.

3.2. Effect of Cur/TBCP2 on CFTR intracellular location

Before testing the Cur/TBCP2 effect, the intracellular distribution of CFTR was examined in F508del-CFTR and normal cells by immunofluorescence and confocal microscopy. The cells were labelled with anti-CFTR antibodies and also with antibodies directed against calreticulin or calnexin which are two quality-control chaperones that bind to misfolded proteins such as the muted CFTR and prevent them from being exported from the ER to the Golgi apparatus for complete glycosylation. The cells were also stained with antibodies directed against ergic-53 because previous observations suggested that F508del-

CFTR was also present in the Endoplasmic reticulum intermediate compartment (ERGIC) [35]. Confocal microscopy images showed that in normal (16HBE14o-) cells, CFTR was located at the periphery of the cells or at the plasma membrane (blue spots) and very few colocations (white spots) were observed with calreticulin or calnexin (Fig. 5). CFTR was logically detected also in ergic-53 en route for its glycosylation in the Golgi. In Σ CFTE29o- cells, the F508del-CFTR was mostly concentrated in the perinuclear region and strongly colocalized with antibodies directed against calnexin and ergic-53 in line with the retention of the mutated protein in the ER/ERGIC compartments of cells expressing the F508del-CFTR (Fig. 5). The retention effect in ERGIC looked stronger in Σ CFTE29o- cells with the presence of many white spots and no blue ones than in CFBE41o- cells. The F508del-CFTR distribution in CFBE41o- looked like that the CFTR in 16HBE14o- cells suggesting localization in the plasma membrane. However, it was reported that the F508del-CFTR was distributed in the cytoplasm of CFBE41o- cells [36]. The CFBE41o- cells are bronchial epithelial cells while the Σ CFTE29o- cells are from tracheal epithelial cells. The intracellular distribution of the F508del-CFTR could vary with the nature of the pulmonary epithelial tissues [37]. Of note, the F508del-CFTR location in the calreticulin compartment looked weak in the CF cell lines.

When Σ CFTE29o- cells were incubated overnight with 220 μ M Cur solubilized with TBCP2, CFTR was not located in the ER/ERGIC compartment near the nucleus as in absence of Cur but its distribution was similar as in the normal 16HBE14o- cells indicating CFTR redistribution in the presence of Cur (Fig. 6). This redistribution after incubation of CFBE41o- cells with the Cur/TBCP2 solution was not as demonstrative as in Σ CFTE29o- cells (Fig. 6). This could be explained by the lower retention of F508del-CFTR in ERGIC of CFBE41o- cells comparatively to Σ CFTE29o- cells (Fig 5). As expected, no modification was observed in normal 16HBE14o- cells in which CFTR was mostly at the plasma membrane

(Fig. 6). The images reconstitution of several z steps as well as sections of the z-step gallery passing through the middle of a representative Σ CFTE29o-cell showed clearly that F508del-CFTR was concentrated at the nucleus periphery in the absence of Cur while it was distributed throughout the cell and at the plasma membrane in the presence of Cur (Fig. 7). These results evidenced that incubation of F508del-CFTR cells with the Cur/TBCP2 solution promoted the relocation of the CFTR at the plasma membrane.

3.3. CFTR current specific activation by Cur/TBCP2 treatment

Whole-cell patch-clamp experiments were carried out to determine the activation of CFTR chloride current after incubation of CF cells with Cur/TBCP2 solution. First, whole-cell patch-clamp experiments were performed in normal (16HBE14o-) and CF (Σ CFTE29o-) cells in the absence of Cur/TBCP2. The measurement of the associated averaged steady-state current-voltage (I/V) relationships revealed the presence of a high chloride induced current (15.3 ± 2.8 pA/pF) in normal cells (Fig. 8Aa and 8B black squares) whereas in CF cells (Fig. 8Ab and 8B, black circles) a weak chloride current (3.1 ± 0.5 pA/pF) was measured at a potential of + 60 mV. In the presence of 10 μ M CFTRinh-172 - a specific CFTR inhibitor - [38] the chloride current was strongly inhibited in 16HBE14o- cells (Fig. 8B, white squares). In contrast the effect was limited in CF cells (Fig. 8B, white circles). The CFTRinh-172 current sensitivity in 16HBE14o- cells (9.3 ± 2.9 pA/pF at a potential of + 60 mV) was 7-fold higher than in Σ CFTE29o- cells (1.3 ± 0.3 pA/pF). The chloride current measured in both cell lines was also inhibited by 20 μ M GlyH-101 - another specific CFTR inhibitor - [39] (data not shown). All together these data demonstrated that chloride currents measured on these different cell lines corresponded to a CFTR chloride current and were in line with those as previously described for these two cell lines [40, 41].

Figure 9 presents characteristic whole-cell currents in Cur/TBCP2-treated Σ CFTE29o- cells in the absence and the presence of 10 μ M CFTRinh-172. CFTR currents were

equivalently blocked on this cell lines by CFTRinh-172 and GlyH-101, the experiments were then conducted with CFTRinh-172 only. Compared to untreated cells, the incubation with Cur/TBCP2 induced a current enhancement that was strongly inhibited in the presence of CFTRinh-172 (Fig. 9A). The CFTRinh-172 current sensitivity was calculated as the difference between currents measured prior and after CFTR-inh 172 exposure, in the same way as done for Fig. 9B. CFTRinh-172 current sensitivity plotted as current-voltage (I-V) relationships revealed that incubation of Σ CFTE29o- cells with Cur/TBCP2 (120 μ M) induced a statistically significant increase in CFTRinh-172 current sensitivity at all membrane potential tested (Fig. 9B). Cur/TBCP2 induced a linear conductance typical of a Cl^- selective current. At +30mV, the current density reached 4.5 ± 1 pA/pF with Cur/TBCP2 *versus* 1 ± 0.6 pA/pF under basal conditions. The observed zero current potential was -39.3 ± 0.1 mV (n= 19) a value that closely matched the chloride reversal potential (-33.8 mV) predicted by the Nernst equation with an internal $[\text{Cl}^-]$ of 42 mM and external $[\text{Cl}^-]$ of 160 mM, suggesting channel selectivity for Cl^- . All together, these results strongly supported that Cur/TBCP2 induced the expression of a Cl^- selective current at the membrane surface of Σ CFTE29o- cells typical of new functional CFTR channels.

2.5. CFTR Cl^- activity assessed by MQAE fluorescence assay

The functional measurement of the CFTR expression in the different airway epithelial cells was also assessed by using the chloride fluorescent probe MQAE [33]. The fluorescence of MQAE is quenched in the presence of chloride or iodide ions in CF cells but its fluorescence is restored in response to $\text{I}^- / \text{NO}_3^-$ substitution and cAMP stimulation after CFTR restoration. Figure 10 presents MQAE fluorescence measurements using I^- , NO_3^- , $\text{NO}_3^- +$ cAMP (8-CPT-cAMP) solution substitution protocol in normal 16HBE14o-, untreated and Cur/TBCP2-treated CFBE41o- cell monolayers. In contrast to 16HBE14o- cells (Fig. 10A),

MQAE fluorescence was not changed in untreated CFBE41o- cells upon I⁻ substitution by NO₃⁻ or was slightly induced by cAMP + NO₃⁻ (amplitude 10.96 ± 2.6) (Fig. 10B).

The incubation of the CFBE41o- cell monolayer with Cur/TBCP2 (165 μM) drastically restored the MQAE fluorescence response to I⁻ / NO₃⁻ substitution (amplitude 24.4 ± 4.4) and particularly in response to cAMP stimulation (amplitude 39.55 ± 6.6) (Fig. 10C). The fluorescence amplitude variation of the fluorescence intensity used as an index of CFTR activity demonstrated a specific restoration of the CFTR activity in CF cells treated with Cur/TBCP2 close to that observed in normal cells (Fig. 10D). Of note, MQAE fluorescence testing was not performed with ΣCFTE29o- cells because the functional restoration of F508del-CFTR was convincing by Patch-clamp measurements (Fig 9). In contrast it was less demonstrative with CFBE41o- cells.

3. Discussion

Several strategies have been explored to improve solubility and stability of Cur in water. For instance, Cur was conjugated to sugars [42, 43], amino acids [44] and polyethylene glycol [45, 46]. Formulations with carriers were also explored [47, 48]. For instance, Cur solubility and activity were improved upon inclusion in cyclodextrin [49] or encapsulation in liposomes [50-52], chitosan/poly(butyl cyanoacrylate) nanoparticles [53, 54], PLGA nanoparticles [55] or methoxy poly(ethylene glycol)-block-polycaprolactone diblock copolymers micelles [56]. Cur effect was enhanced in CF mice upon oral administration of Cur encapsulated in biodegradable nanoparticles made of an oil-in-water emulsion mixture of poly (lactic-co-glycolic acid) (PLGA) and poly(vinyl alcohol) [57]. ABA copolymers comprising poly(ethylene oxide) (PEO) block as the hydrophilic moiety and poly(propylene oxide) (PPO) block as hydrophobic moiety forming core-shell micelles were also used. However, PEO polymers due either to the polymer itself or a side product formed during its synthesis can generate various unfavourable effects [58]. Polyoxazoline was proposed as an

alternative to PEO [15, 59] and a family of poly(2-methyl-2-oxazoline-b-tetrahydrofuran-b-2-methyl-2-oxazoline) (MeOx-THF-MeOx) neutral amphiphilic triblock copolymers (TBCP) have been synthesized [20]. Our Cur/TBCP2 formulation (17 wt.%) was obtained by solubilisation of Cur (0.5 mg) in a water solution of micelles of TBCP2 copolymer (2 mg/ml; 2400 g.mol⁻¹). Comparatively, the preparation of 1 mg/ml curcumin with 10 mg/ml of 1,2-dimyristoyl-sn-glycero-3-phosphocholine and 1,2-dimyristoyl-sn-glycero-3-phospho-rac-(1-glycerol) liposomes was described [52]; the preparation of 14% wt.% and 7.6% wt.% curcumin formulations were reported with methoxy poly(ethylene glycol)-block-polycaprolactone diblock copolymers micelles [56] and poly (lactic-co-glycolic acid) (PLGA) and poly(vinyl alcohol) nanoparticles [57], respectively. With another highly hydrophobic drug, the preparation of a 45 wt.% Paclitaxel formulation was successfully achieved with the MeOx₃₇-b-BuOx₂₃-b-MeOx₃₇, a triblock copolymer of 10 000 g.mol⁻¹ [15]. The resulting Cur/TBCP2 solution formed nanoparticles of ~250 nm while a 0.2% TBCP2 in water formed micelles of ~16 nm. The large size of TBCP2/Cur nanoparticles could result from the coalescence of TBCP2 micelles around Cur aggregates. Here, we show that TBCP2 allowed Cur internalization by human airway epithelial cell lines. Yet, the penetration of Cur was lower than when Cur was solubilized in DMSO which allows Cur diffusion through the plasma membrane. The uptake mechanism of Cur in TBCP2 micelles is not yet determined but it could process *via* a transient destabilization of the plasma membrane by TBCP2. Indeed, Rho-TBCP does not enter deeply into the cytoplasm of the tracheal epithelial cells while Cur enters the cells. This suggests that once bound to the plasma membrane, the copolymer would destabilize the lipid bilayer inducing Cur diffusion through the plasma membrane. The copolymer that is more internalized in the bronchial epithelial cells would in addition induce the formation of transient pores in endocytic vesicles allowing Cur diffusion in the cytoplasm. This latter hypothesis is supported by data showing that a neutral

416 amphiphilic diblock copolymer - a poly(ethylene oxide-b-4-vinylpyridine) - was able to form
417 transient pores in a lipid artificial membrane allowing the passage of a plasmid DNA of ~5
418 kbp across the model membrane [60].

419 Here, we have evaluated the biofunctionality of curcumin after solubilisation by
420 TBCP2. For this purpose, we have tested whether the Cur/TBCP2 formulation could restore
421 the expression of a functional CFTR protein at the surface of F508del-CFTR cells. Indeed,
422 F508del-CFTR is not expressed at the plasma membrane because it is retained in the
423 ER/ERGIC compartment by the quality control system involving calreticulin and/or calnexin
424 [61]. Our immunofluorescence analyses performed on two F508del-CFTR cell lines verified
425 that F508del-CFTR colocalizes with calnexin and calreticulin. They confirmed also that
426 F508del-CFTR colocalizes with ERGIC-53, a protein specific of the ERGIC compartment as
427 previously reported [35]. p58/ERGIC-53 is a calcium-dependent lectin recognizing mannose
428 that cycles between the ER and the Golgi apparatus and functions as a cargo receptor for a
429 subset of soluble glycoproteins exported from the ER. The lectin domains of ERGIC-53 and
430 calnexin are structurally similar. F508del-CFTR is not exported to the Golgi for maturation
431 but is driven to the cytosolic ubiquitin-proteasome machinery for degradation. It has been
432 shown that ERGIC compartment accumulates proteins on the way for degradation as the
433 precursor of human asialoglycoprotein receptor H2a and free heavy chains of murine class I
434 major histocompatibility complex (MHC) [62]. Significant amounts of various ER resident
435 proteins have been detected in ERGIC indicating that a leak of calnexin from ER into ERGIC
436 might occur [63]. Thus, calnexin could be transiently present in ERGIC which could be a site
437 for the concentration and retrotranslocation of proteins that are transported to the cytosol.
438 That explains colocalizations between F508del-CFTR and p58/ERGIC-53 observed in the
439 F508del-CFTR cell lines.

Cur was shown to partially compensate the ER retention of the defective CFTR protein in appropriate cell lines and in F508del-CFTR mice [26, 27]. We demonstrate that incubation of F508del-CFTR cell lines with Cur/TBCP2 induces the relocation of the mutated CFTR protein at the plasma membrane. This effect resulted likely from the better cellular penetration of Cur upon solubilization in the TBCP2 solution. The functionality of CFTR chloride channels in the apical membrane of CF cells after incubation with Cur/TBCP2 is clearly revealed by whole-cell patch-clamp experiments and MQAE fluorescence assay. In the former, the specificity of the activation of CFTR currents is provided by the use of CFTRinh-172 or GlyH-101, two specific CFTR inhibitors that both induce strong inhibition of restored currents in CF cells treated with Cur/TBCP2. CFTRinh-172 is a well-known and widely used specific inhibitor of CFTR channels, without affecting other chloride conductance such as the Ca^{2+} -dependent Cl^- conductance (CaCC) or the volume-sensitive outwardly rectifying Cl^- conductance (VSORC). GlyH-101 appears less specific as it inhibits both VSORC and CaCC at concentrations close to those used to inhibit CFTR conductance [64]. Therefore, our calculated CFTRinh-172 current sensitivity was due to CFTR current. The fluorescence assay with the chloride sensitive fluorescent probe MQAE which is quenched in the presence chloride or iodide ions is drastically restored in CF cells after Cur/TBCP2 treatment in response to $\text{I}^- / \text{NO}_3^-$ substitution and cAMP stimulation. All together, these results provide strong evidences that the CFTR relocation leads to a functional chloride channel.

The exact mechanism by which Cur allows the F508del-CFTR relocation is not fully understood. Cur is an inhibitor of the sarcoplasmic/endoplasmic reticulum Ca^{2+} -ATPase (SERCA) involved in the translocation of calcium from the cytosol to the sarcoplasmic reticulum lumen [65-67]. The lectin-like recognition of calreticulin, calnexin and ERGIC-53 that binds to the terminal glucose of the N-oligosaccharide structure of misfolded

glycoproteins is calcium-dependent. SERCA inhibition by Cur *via* the modulation of sarcoplasmic reticulum calcium content could prevent retention of F508del-CFTR in the ER/ERGIC favouring its delivery in the Golgi apparatus for maturation and exocytosis to the plasma membrane. It has also been reported that the trafficking of CFTR to the plasma membrane involves the keratin 8/keratin 18 network of intermediate filaments [68, 69]. An important remodelling in the keratin 18 network has been indeed observed in Cur-treated cells with an increase in the phosphorylation of K18 Ser52 in a calcium-independent manner [70]. This reorganization reduces the intracellular trafficking of organelles mediated by the intermediate filaments and the turnover of some membrane proteins as CFTR. Inhibitions by Cur of the ER retention of F508del-CFTR and its turnover from the plasma membrane could contribute to the restoration of a functional CFTR in the plasma membrane. Curcumin was proposed to treat Cystic Fibrosis but its multiple therapeutic effects as well as the absence of fully understanding mechanism (s) of action delay its FDA agreement. For this reason, curcumin is often named as pan-assay interference (PAIN) compound, classes of compounds that can interfere with bioassays *via* a number of different mechanisms [71].

4. Conclusion

We demonstrated the great potential of the MeOx₆-THF₁₉-MeOx₆ copolymer to solubilize a very insoluble molecule such as curcumin. In a cystic fibrosis context, we showed that curcumin better penetrated normal and F508del-CFTR human airway epithelial cells and promoted a functional expression of the mutated CFTR protein in the plasma membrane of the CF cells. More generally, MeOx_n-THF_m-MeOx_n copolymers could help solubilisation of other water insoluble drugs or cosmetic ingredients contributing to their better applications.

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Figure 1: (A) Chemical structure of the amphiphilic triblock copolymer MeOx₆-THF₁₉-MeOx₆ (TBCP2). (B) Cur solubilisation in a TBCP2 aqueous solution. Two mg Cur was added to 5 ml of a 0.2% TBCP2 solution in water (+TBCP2) or in 5 ml water (-TBCP2). (C) ¹H NMR spectrum of TBCP2. (D) Size of TBCP2 micelles and Cur/TBCP2 formulation.

Figure 2: TBCP2 allows Cur penetration in the cells. Σ CFTE290-, 16HBE140- and CFBE410- cells were incubated at 37°C in the presence of (A) 220 μ M Cur solubilized in TBCP2, (B) supernatant of 220 μ M Cur in PBS or (C) TBCP2 without Cur. (D) as control, the cells were incubated without any Cur/TBCP2 formulation. After 2 h, the cell fluorescence was analyzed by confocal laser scanning microscope (λ_{ex} : 488 nm; λ_{em} : 520 nm). Bar scale: 20 μ m.

Figure 3: Uptake of Cur. (◆, ◇) ΣCFTE29o-, (■, □) 16HBE14o- and (▲, Δ) CFBE41o- cells were incubated for 2 h at 37°C in the presence of various dilutions of the Cur/TBCP2 formulation. The mean fluorescence intensity (MFI) of the cells was measured by flow cytometry (λ_{ex} : 488 nm; λ_{em} : 530 ± 30 nm) before (black symbols) and after (white symbols) treatment with trypan blue. The fluorescence intensity is expressed as MFI value of 10,000 cells.

Figure 4: Curcumin and copolymer cellular distribution. Σ CFTE29o-, 16HBE14o- and CFBE41o- cells (1.4×10^4) were seeded on glass coverslips in 2 cm^2 well of a 4-well plastic culture plate. Two days after, the cells were incubated at 37°C in the presence of $40 \mu\text{M}$ curcumin solubilised with Rho-TBCP. Upon 4h incubation, cells were washed with PBS coverslips were mounted in Vectashield and analysed by confocal laser scanning microscopy. The fluorescence of curcumin and rhodamine were measured at 520 nm upon excitation at 488 nm and 560 nm upon excitation at 543 nm, respectively. Scale bar: $20 \mu\text{m}$.

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Figure 5: Confocal microscopy images of CFTR intracellular location. Σ CFTE29o-, 16HBE14o- and CFBE41o- cells were labelled with mouse anti-hCFTR antibodies and either with anti-calreticulin anti-calnexin or anti-ERGIC-53 antibodies. Anti-hCFTR antibodies were revealed with Cy5 sheep anti-mouse antibodies (blue) and the other antibodies with Alexa Fluor 568 secondary antibodies (red). Scale bar: 20 μ m.

Figure 6: Effect of Cur/TBCP2 treatment on the F508del-CFTR location. Σ CFTE29o-, CFBE41o- and 16HBE14o- cells were cultured for 16 h in the absence (without Cur) or the presence (with Cur) of 220 μ M Cur/TBCP2. The cells were stained with mouse anti-hCFTR antibody followed by Cy5-sheep anti-mouse antibodies. The fluorescence of the cells was analysed by confocal microscopy at 660 nm upon excitation at 633 nm (Cy5). Scale bar: 20 μ m.

Figure 7: Images reconstitution of several z steps (A, C) and z-step gallery (B, D) after (A, B) and before (C, D) Cur/TBCP2 treatment of Σ CFTE29o- cells.

Figure 8: Patch-clamp characterization of CFTR current in normal and CF epithelial cells. (A) Representative whole-cell Cl^- current recordings in (a) 16HBE14o- and (b) Σ CFTE29o- cultured cells under control conditions or (c and d) after addition of the selective CFTR-inhibitor CFTRinh-172 (10 μM) for 10 min into the bath solution. (B) Plots of averaged steady-state current-voltage relationships (I/V curves) of Cl^- currents in the absence (black squares: 16HBE14o- cells, n=5; black circles: Σ CFTE29o- cells, n=9) or after addition of 10 μM CFTRinh-172 for 10 min (white symbols). All results were expressed as mean $\pm$ SEM.

Figure 9: Patch-clamp measurements of Cur/TBCP2-induced activation of CFTRinh-172 sensitive current in Σ CFTE29o- cells. (A) Representative whole cell CFTR current traces recorded after cell culture (a) in basal conditions or after 16 h treatment with (b) Cur/TBCP2 (120 μ M) or (c) after addition of the selective CFTR-inhibitor CFTRinh-172 (10 μ M) for 10 min into the bath solution in cells treated with Cur/TBCP2. (B) Plots of averaged steady-state current-voltage relationships (I/V curves) of CFTRinh-172 sensitive currents. I/V curves on cells before treatment (black squares, n= 8), and 16 h treatment with Cur/TBCP2 (black triangles, n= 19). Steady-state current amplitude was measured at the end of the pulse and normalized to the cell capacitance. All results were expressed as mean $\pm$ SEM. Statistical significance was assessed using Mann-Whitney non-parametric test, and the significance level was established at $p < 0.05$ between CFTRinh-172 sensitive currents measured with Cur/TBCP2.

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Figure 10: Cur/TBCP2 treatment influence on the CFTR-mediated anion transport on different epithelial cell monolayers measured by MQAE fluorescence. CFBE41o⁻ and 16HBE14o⁻ cells loaded with MQAE were sequentially perfused with I⁻, NO₃⁻, NO₃⁻ with cAMP and again with I⁻ buffer solutions. Representative MQAE fluorescence intensity was plotted as a function of time in (A) 16HBE14o⁻, (B) untreated CFBE41o⁻ cells or (C) treated CFBE41o⁻ cells for 16 h with Cur/TBCP2 (165 μM). (D) Histograms of fluorescence amplitude variation (Δ fluorescence) in cell monolayer without or with Cur/TBCP2 treatment, in the presence of NO₃⁻ and NO₃⁻ + cAMP (grey filled histograms) or NaI⁻ (white histograms) expressed as the difference between the maximal fluorescence intensity measured at steady state before and after the change of the respective solutions. Data are means ± SEM. Statistical significance was assessed using a Student's *t*-test, and the significance level was established at *p*<0.05 (*: *p*<0.05; **: *p*<0.01; ***: *p*<0.00.1). ns: non-significant.

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solubilization Cell penetration

CFTR relocalisation

Chloride transport restoration

Graphical Abstract