



HAL
open science

Sequential delivery of synergistic drugs by silica nanocarriers for enhanced tumour treatment

Albane Birault, Simon Giret, Christophe Théron, Audrey Gallud, Afitz da Silva, Denis Durand, Christophe Nguyen, Nadir Bettache, Magali Gary-Bobo, John Bartlett, et al.

► To cite this version:

Albane Birault, Simon Giret, Christophe Théron, Audrey Gallud, Afitz da Silva, et al.. Sequential delivery of synergistic drugs by silica nanocarriers for enhanced tumour treatment. *Journal of materials chemistry B*, 2020, 8 (7), pp.1472-1480. 10.1039/c9tb02225b . hal-03096456

HAL Id: hal-03096456

<https://hal.science/hal-03096456v1>

Submitted on 5 Jan 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Sequential Delivery of Synergistic Drugs by Silica Nanocarriers for Enhanced Tumour Treatment

Albane Birault,^a Simon Giret,^a Christophe Théron,^a Audrey Gallud,^b Afitz Da Silva,^b Denis Durand,^b Christophe Nguyen,^b Bettache Nadir,^b Magali Gary-Bobo,^b John R. Bartlett,^{c,d} Michel Wong Chi Man,^a Carole Carcel^{a*}

^aICGM, Univ Montpellier, CNRS, ENSCM, Montpellier, France

^bInstitut des Biomolécules Max Mousseron, UMR5247, Université de Montpellier, CNRS, ENSCM, Faculté de Pharmacie, 15 Avenue Charles Flahault, 34093, Montpellier, Cedex 05, France

^cUniversity of the Sunshine Coast, 90 Sippy Downs Drive, Sippy Downs QLD 4556, Australia

^dWestern Sydney University, Locked Bag 1797 Penrith, NSW, 2751, Australia

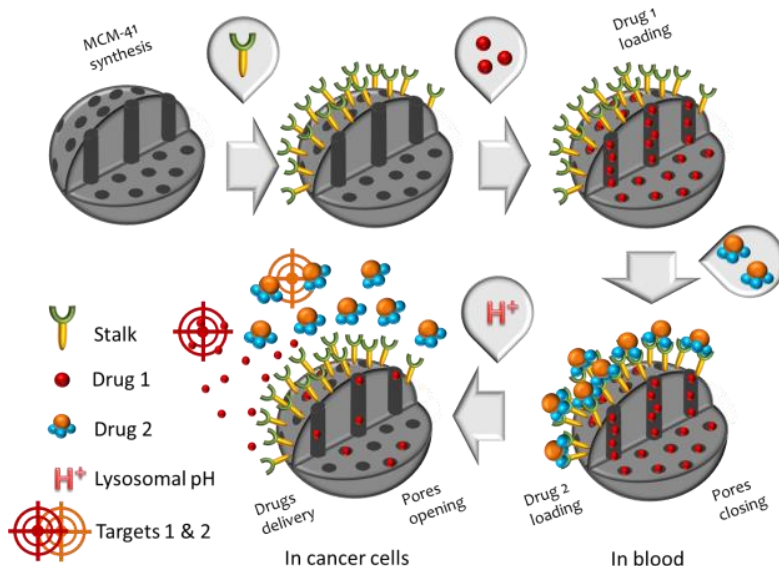
Herein hybrid silica nanoparticles have been engineered to direct the sequential delivery of multiple chemotherapeutic drugs in response to external stimuli such as variations in pH. The nanocarriers consist of conventional MCM-41-type nanoparticles, which have been functionalised with an organic ligand (or stalk) grafted onto the external surface. The stalk is designed to “recognise” a complementary molecule, which serves as a “cap” to block the pores of the nanoparticles. First, camptothecin is introduced into the pores by diffusion prior to capping the pore apertures *via* molecular recognition. The cap, which is a derivative of 5-fluorouracil, serves as a second cytotoxic drug for synergistic chemotherapy. *In vitro* tests revealed that negligible release of the drugs occurred at pH 7.4, thus avoiding toxic side effects in the blood stream. In contrast, the stalk/cap complex is destabilised within the endolysosomal compartment (pH 5.5) of cancer cells, where release of the drugs was demonstrated. Furthermore, this environmentally responsive system exhibited a synergistic effect of the two drugs, where the pH-triggered release of the cytotoxic cap followed by diffusion-controlled release of the drug cargo within the pores led to essentially complete elimination of breast cancer cells.

Introduction

Combinatorial chemotherapy is widely employed as a primary treatment approach in cancer therapy to overcome the easily-developed defence of cancer cells against a single drug.^{1, 2} However, conventional therapies involving the use of multiple free drugs often lead to well-known severe toxic side effects.^{2, 3} One of most effective approaches for overcoming such limitations is to load multiple therapeutic agents that exploit different anti-tumour mechanisms into a single effective nanocarrier with no drug leakage before reaching the target. Such smart nanocarriers could then increase the therapeutic efficiency and reduce toxic side effects by improving the bio-accessibility of drugs.⁴⁻⁸ In this sense, mesoporous silica nanoparticles (MSNPs) have been widely used, due to their versatility in forming the basis of stimuli-responsive drug delivery systems.⁹⁻¹³ Notably, pH-responsive MSNPs provide the advantage of being endogenously stimulated due to the difference in pH between cellular compartments.¹⁴⁻¹⁶ We recently reported¹⁷ the use of molecular recognition *via* H-bonding to design smart silica nanoparticles with a pH-activated cap, which exhibited slow release kinetics under slightly acidic conditions and good efficacy for cancer cells elimination.

In order to avoid the well-known phenomenon of anti-cancer drug resistance and to optimize this earlier nanosystem, the current work explored the design of a multi-drug nanoplatfrom for combined chemotherapy. Our strategy, outlined in **Scheme 1**, involves coupling silica nanocarriers embodying molecular recognition sites with an active cap to enhance the anti-tumour activity. After loading the functionalised nanoparticles with camptothecin (CPT, **Drug 1**), the pores are closed to avoid premature release under physiological conditions within the bloodstream (pH 7.4) using a 5-fluorouracil (5-FU) derivative (**Drug 2**), which can create stable H-bonds with **Stalk** located on the nanoparticles' surfaces. The opening of the pores is then triggered at the acidic pH within lysosomes (pH 4.5–5.5), to deliver sequentially the two drugs within the cancer cells because of **Stalk** protonation followed by disruption of the **Stalk-Cap** bonds.

As already mentioned, there are a plethora of studies reporting MSNPs efficacy for cancer therapy applications.^{18, 19} Although some of these have described the use of drugs as pore-blockers to form promising nanocarriers for combination therapy,²⁰⁻²² to the best of our knowledge, this is the first time that a pH-responsive nanocarriers has been employed to exploit the synergistic toxicity of combining CPT and 5-FU for anti-cancer drug resistance applications.



Scheme 1. Conceptual combinational drug delivery MSNP nanocarriers with Drug 1 (camptothecin) encapsulated inside the pores and Drug 2 (5-FU derivative) as the capping agent

To develop such systems, two separate syntheses are required: 1) the functionalisation of appropriate silica nanocarriers such as MCM-41, by **Stalk** grafting, to provide selective H-bonding recognition sites on the external surface; and 2) the synthesis of an antitumor derivative, with a complementary molecular recognition pattern, which, by pairing with the corresponding **Stalks**, would then cap the pores.

Organosilylated triazine derivatives, bearing donor-acceptor-donor recognition sites, were used as **Stalks** condensed onto the surface of the nanoparticles. The **Stalk** employed in this study is tetrasilylated, which enhances the functionality of the platform by providing additional degrees of freedom and binding sites for grafting the ligand onto the surface of the nanoparticles, compared to our previously described bisilylated system.¹⁷ The additional conformational flexibility of the tetrasilylated **Stalk** would also be expected to enable the amine sites on the ligand to orient correctly to form the desired H-bonds with the **Cap**. As drug candidates, we chose for the pore-blocker a commonly used anticancer drug, 5-fluorouracil (5-FU), which exhibits complementary molecular recognition properties to **Stalk**. Previously,²³ we have demonstrated that a derivative such as **Cap** (Figure 1) with a three-fold H-bonding pattern favours stable complexation at neutral pH in bulk hybrid silica materials. In contrast, the complex begins to dissociate at lower pH after triazine unit protonation, leading to the breakdown of H-bonds initially involved in stabilization of the capping complex, thus promoting removal of **Cap** at pH 5.5 and below and its associated release. The biological activity of **Cap** was previously evaluated²³ by cytotoxic assay on MCF-7 human breast cancer cells, showing a relatively high efficiency (43% of breast cancer cell death at 50 μ M) for this molecule as an anticancer drug compared with the parent 5-FU (68% of cell death at 50 μ M).

To investigate their viability and the combination therapeutic effect, we evaluated the cytotoxicity of these MSNPs towards human breast cancer cells (MCF-7). In a first study, the functionalised nanoparticles were loaded with fluorescent dyes, such as propidium iodide (PI) or rhodamine-B (RhB), to validate the opening of the pores in weakly-acidic lysosomal media and to assess the internalization of the MSNPs in the cells. In a second study, the MSNPs were loaded with CPT and the pores capped with 5-FU derivative **Cap** to examine the overall cytotoxicity by *in vitro* studies. CPT destroys cancer cells *via* a different biological pathway than 5-FU^{24, 25} and the two drugs have been previously shown to demonstrate a strong synergistic effect when used together to treat cancer cells.²⁶ The approach outlined in this study enables two cytotoxic drugs to be delivered autonomously and sequentially, which is typically the manner in which injection- or drip-based combination drug therapies are administered to maximise the efficacy of such treatment. Under *in-vitro* conditions, the multi-drug payload is retained under normal physiological conditions and then released autonomously at the disease site, demonstrating nearly complete cell apoptosis within 72 h.

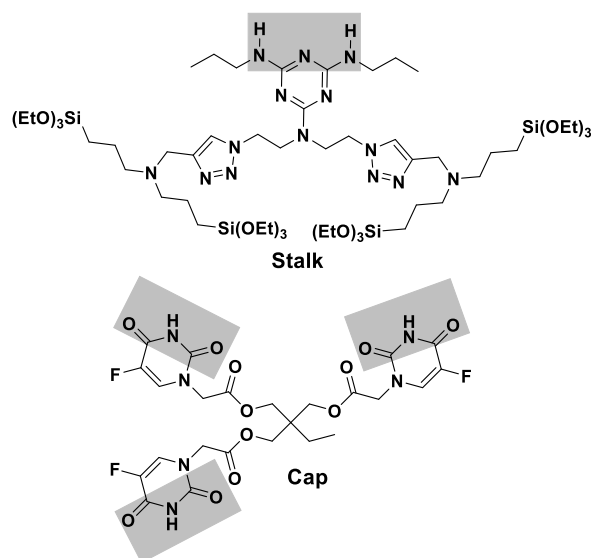


Figure 1. Structures of Stalk precursor and Cap

Experimental Section

General

5-fluorouracil, thionyl chloride, triethylamine and 1,1,1-tri(hydroxymethyl)propane were purchased from Sigma, USA and were used without purification. Solvents were dried by employing a MB SPS-800 apparatus.

Characterization

FTIR spectra were obtained on a Perkin Elmer FT-IR Spectrum BX spectrometer. CPMAAS solid-state NMR spectra were obtained using a Bruker FT-AM 400 spectrometer and ^1H - and ^{13}C -NMR spectra were obtained using a Bruker AC-400 spectrometer, with CDCl_3 or DMSO-d_6 as solvent and tetramethylsilane (TMS) as a reference. N_2 adsorption/desorption isotherms were obtained using a Micromeritics ASAP 2010 instrument at 77.15 K. Specific surface areas were calculated using the BET transform directly from the isotherms, using 0.162 nm^2 as the cross-sectional area of N_2 . Electron micrographs were obtained with a JEOL 1200 EXII microscope for TEM and with a Hitachi S4800 30kV apparatus for the SEM data. ^{29}Si solid-state NMR spectra were obtained using a Bruker DSSX 300 MHz spectrometer. Spectra were recorded using 4000 scans, with a relaxation delay of 3 sec, pulse duration of 6 μsec and acquisition time of 0.04 sec.

Thermogravimetric analyses (TGA) were performed on a Netzsch TG 209 C apparatus employing a heating rate of $10 \text{ }^\circ\text{C min}^{-1}$ under an air flow of 20 mL min^{-1} up to 585°C . The small and wide-angle X-ray scattering (SWAXS) experiments were conducted using a Guinier Mering set up with a 2D image plate detector.

Precursor Synthesis

Synthesis of 5-FUA

5-FUA was prepared according to a previously reported method.²⁷ Yield: 54%; m.p. $274\text{--}276 \text{ }^\circ\text{C}$;

^1H NMR (400 MHz, DMSO-d_6): $\delta=11.93$ (s, 1H), 8.09 (d, 1H), 4.37 ppm (s, 2H); ^{13}C NMR (100 MHz, DMSO-d_6 , δ): $\delta=168.6, 158.2, 150.8, 140.1, 128.2, 53.4$; HRMS (ESI) m/z: $[\text{M} + \text{H}]^+$ calcd for $\text{C}_6\text{H}_6\text{N}_2\text{O}_4\text{F}$: 189.0312; found: 189.0313.

Synthesis of Cap (Figure 3)

The synthesis of **Cap** was adapted from a previously reported procedure.²² **5-FUA** (1.1 g, 5.85 mmol) and triethylamine (126 μL , 0.904 mmol) were dissolved in dry THF (44 mL) and freshly distilled SOCl_2 (8.4 mL) at $20 \text{ }^\circ\text{C}$ under N_2 . After 3.5 h, excess SOCl_2 and THF were removed under vacuum and the residue was re-dissolved in dry THF (42 mL). 1,1,1-tri(hydroxymethyl)propane (80 mg, 0.60 mmol) was added and the mixture was stirred at $20 \text{ }^\circ\text{C}$ for 48 h. The solvent was then evaporated, and the residue was purified by column chromatography ($\text{CH}_2\text{Cl}_2/\text{MeOH}$: 98:2 90:10). Yield: 72%; m.p. decomposition after $180 \text{ }^\circ\text{C}$;

^1H NMR (400 MHz, DMSO-d_6): $\delta=12.00$ (s, 3H), 8.04 (d, 3H), 4.50 (s, 6H), 4.05 (s, 6H), 1.36 (q, 2H), 0.79 ppm (t, 3H); ^{13}C NMR (100 MHz, DMSO-d_6): $\delta=20.9, 34.7, 48.7, 63.4, 130.1, 138.1, 140.5, 149.6, 157.2, 167.6$ ppm; HRMS (ESI) m/z: $[\text{M} + \text{H}]^+$ calcd for $\text{C}_{24}\text{H}_{24}\text{N}_6\text{O}_{12}\text{F}_3$: 645.1404; found: 645.1412.

Synthesis of Stalk (Figure 2)

The **Stalk** was prepared according to a previously reported procedure.²⁸ Firstly, compound **1** was added dropwise (due to its potentially-explosive nature, care should be taken) to a solution of cyanuric chloride in THF in the presence of N,N-diisopropylethylamine (DIPEA) at 0 °C, to afford **2** (66% yield). Compound **2** was then dissolved in THF with 10 equivalents of propylamine under reflux to give **3** (90% yield). Finally, the tetrasilylated **Stalk** was obtained *via* a CuAAC Click coupling reaction between **3** and **4**. The reagents were introduced into a microwave oven in the presence of the CuBr(PPh₃)₃ catalyst and Et₃N in THF at 100 °C for 20 minutes to yield the **Stalk (Figure S1)** in quantitative yield.

¹H NMR (400 MHz, CDCl₃, **Figure S1**): δ = 0.56 (t, 8H), 0.95 (t, 6H), 1.22 (t, 36H), 1.58 (m, 12H), 2.39 (t, 8H), 3.29 (q, 4H), 3.64 (m, 8H), 3.80 (q, 24H), 4.47 (t, 4H), and 7.32 (s, 2H). ¹³C NMR (100 MHz, CDCl₃): δ = 7.86 (s), 11.50 (s), 18.28 (s), 20.26 (s), 23.05 (s), 42.47 (s), 45.90 (s), 48.16 (s), 48.60 (s), 56.46 (s), 58.29 (s), 123.11 (s), 128.51 (s), and 132.01 (s). HRMS (ESI) m/z: [M + H]⁺ calcd for C₅₅H₁₁₅N₁₄O₁₂Si₄ (MH): 1275.7896; found: 1275.7910.

Nanoparticles preparation (Scheme 2)

Synthesis of NP

Cetyltrimethylammonium bromide (CTAB; 315 mg, 8.6x10⁻¹ mmol), was dissolved in Milli Q H₂O (150 mL, 8.3x10³ mmol), and the pH was adjusted to ~ 12 by addition of NaOH solution (2 M, 1.1 mL). After heating the resulting solution to 80 °C, tetraethylorthosilicate (TEOS; 1.4 mL, 6.3 mmol) was added dropwise and the reaction was continued for 2 h to yield MCM-41 nanoparticles. Finally, the nanoparticles were collected after centrifugation and washing with EtOH. ²⁹Si CPMAS solid-state NMR: δ=-99.2, -108.9 ppm

Synthesis of NP1

Stalk (80 mg, 0.10 mmol) was added to a suspension of MCM-41 (80 mg) (pre-heated overnight at 90 °C to remove traces of H₂O and to activate Si-OH functions) in dry toluene (10 mL) under an inert atmosphere. The suspension was then heated at 80 °C for 24 h, prior to centrifuging the nanoparticles and washing them twice with ethanol and then five times with water. ²⁹Si CPMAS NMR: δ=-57.4, -100.1, -108.9 ppm.

Synthesis of NP2

The CTAB surfactant was removed by washing with a solution of HCl 37 wt% (1 mL) in ethanol (80 mL) under reflux overnight. The resulting **NP2** nanoparticles were washed twice with EtOH.

Synthesis of NP3

The amino groups of the triazine derivative on the surface of **NP2** were deprotonated by washing the nanoparticles with freshly distilled Et₃N (1 mL, 7.2 mmol) in H₂O (10 mL, 5.6x10² mmol) for 48 h at 70 °C. The resulting **NP3** nanoparticles were then washed two times with H₂O and three times with EtOH.

Loading, capping and release

NP3 nanoparticles were suspended in a solution of cargo molecules as described below. The suspension was then sonicated for 20 min and stirred for 18 h to promote filling of the pores with the cargo molecules. To ensure retention of the drug, the pore apertures were capped by addition of **Cap** (3 mg; 4.7x10⁻³ mmol) and the suspension was stirred for additional 48 h. After centrifugation, the particles were washed with DMSO around 5-6 times following by washing with water and EtOH.

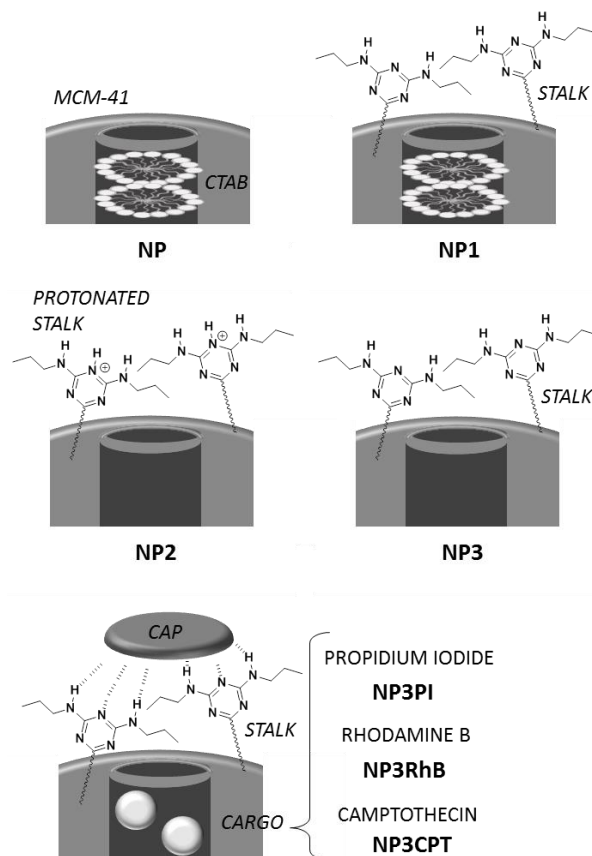
NP3PI (5 mM): PI (45 mg, 6.7x10⁻² mmol) in water (13.5 mL), and nanoparticles (90 mg);

NP3RhB (5 mM): RhB (32 mg, 6.7x10⁻² mmol) in water (13.5 mL), and nanoparticles (90 mg);

NP3CPT (3 mM): CPT (15 mg, 4.36.7x10⁻² mmol) in DMSO (15 mL), and nanoparticles (90 mg).

The CPT loadings obtained under the conditions used to impregnate the **NP3** nanoparticles were 4 wt%, on the basis of visible absorbance spectroscopy studies.

The release of the CPT from **NP3CPT** mesopores was evaluated by UV-Vis absorbance at 380 nm at either pH 2, 5.5 or 7.4 for different periods of time over a 24 h period. For these studies, 3 mg of **NP3CPT** was dispersed into 1 mL of aqueous solution at the defined pH and then 100 μL aliquots of the nanoparticle dispersion were poured into Eppendorf tubes. The tubes were stirred during release of the drug and were then centrifuged (15x10³ min⁻¹, 15 min). The resulting supernatant was placed into a microreader plate and the concentration of drug was determined from an appropriate calibration curve.



Scheme 2. Synthesis of capped nanoparticles

In vitro studies

Cell culture conditions

Human breast cancer cells (MCF-7) were purchased from ATCC (American Type Culture Collection, Manassas, VA). MCF-7 cells were cultured in DMEM-F12 culture medium supplemented with 10 % foetal bovine serum and 50 $\mu\text{g}\cdot\text{mL}^{-1}$ gentamycin. These cells were allowed to grow in a humidified atmosphere at 37 °C under 5 % CO_2 .

Cytotoxicity

MCF-7 cells were seeded into 96-well plates at 10^4 cells per well in 200 μL culture medium and allowed to grow for 24 h. The three batches of nanoparticles (**NP**, **NP3RhB**, and **NP3CPT**) were dispersed in ethanol at a concentration of 10^{-2} M and sonicated in an ultrasonic bath until completely dispersed. Then, cells were incubated for 72 h with different nanoparticles concentrations (from 1 to 100 $\mu\text{g}\cdot\text{mL}^{-1}$). For the **NP3CPT** batch, the experiment was also carried out for shorter lengths of time (6, 16 and 24h). At the end of the incubation time, a MTT assay was performed to evaluate the toxicity. Briefly, cells were incubated for 4 h with 0.5 $\text{mg}\cdot\text{mL}^{-1}$ of MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazoliumbromide; Promega) in culture media. The MTT/media solution was then removed, and the precipitated crystals were dissolved in EtOH/DMSO (1:1). The solution absorbance was read at 540 nm.

Confocal imaging

MCF-7 cells were seeded one day prior to nanoparticle exposure at 10^6 cells/ cm^2 in glass-bottomed culture dishes from Ibidi Biovalley®. Cells were then exposed for 20 h to 40 $\mu\text{g}\cdot\text{mL}^{-1}$ of NPs. Prior to imaging, the cells were stained with 50 nM LysoTracker Green DND-26 for 30 min and 5 $\mu\text{g}\cdot\text{mL}^{-1}$ Hoechst 33342 was added during the last 10 min of incubation. Confocal images were acquired on a Zeiss Axio Observer confocal microscope equipped with an oil-immersion Plan-Apochromat 63x/1.40 objective.

Flow cytometry

MCF-7 cells were seeded into a 6-well plate (Nunc; 10^6 cells/well) and allowed to grow for 24 h. The cells were harvested at 5, 18 and 22 h following NP exposure (40 $\mu\text{g}\cdot\text{mL}^{-1}$) and re-suspended in DMEM-F12 phenol red-free medium. Dead cells were counterstained with 4',6-diamidino-2-phenylindole (DAPI, 0.5 $\mu\text{g}\cdot\text{mL}^{-1}$). The percentage of

positive living cells for NP uptake was determined on a FACS Canto II flow cytometer. The data were analysed with Win MDI software v2.8.

Results and discussion

Preparation of precursors

The approach used to prepare the **Stalk** (Figures 2 and S1) was based on a previously reported method²⁸ (Figure 2). Similarly, the **Cap** was prepared according to an optimized protocol based on a previously reported synthesis.²³ The reaction was performed in homogeneous medium and employed an eco-friendly room-temperature approach for the second and third steps, which led to an increased yield and a reduction in the overall reaction time (Figure 3).

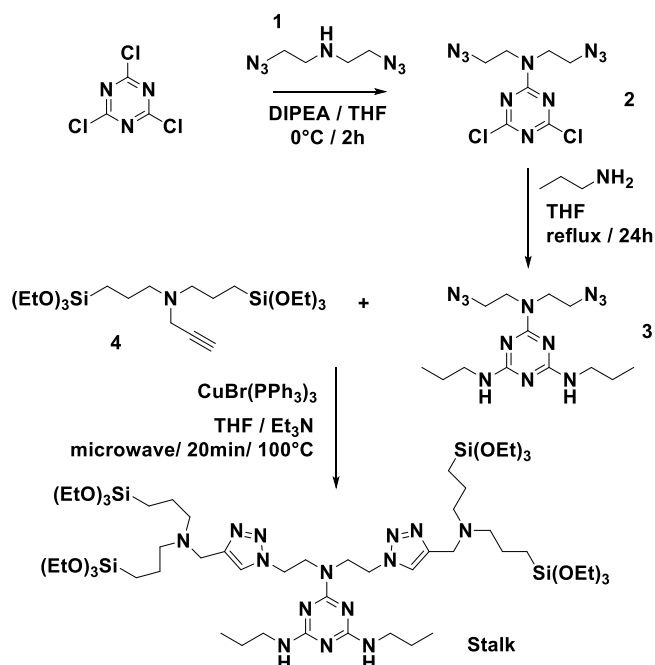


Figure 2. Synthesis of **Stalk**

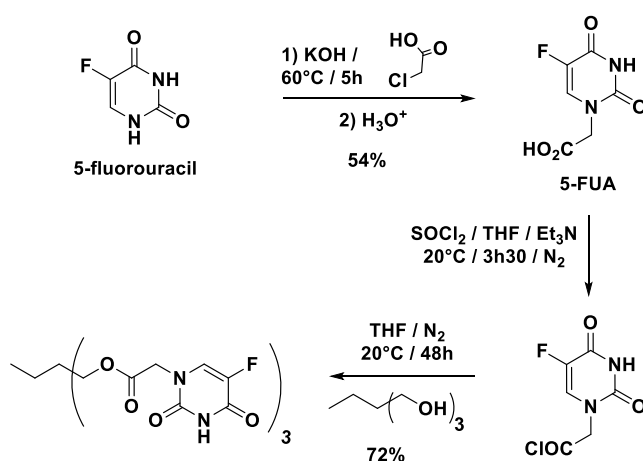


Figure 3. Optimized synthesis of **Cap**

Nanoparticles synthesis, functionalisation, loading and capping

MCM-41 nanoparticles (**NP**) were prepared by classical sol-gel processing in alkaline aqueous solution. CTAB used to template the compact hexagonal porosity (Figure S2) in the MSNPs was retained in the pores, and the **Stalk** was then covalently grafted onto the MSNP surface by condensation. After washing with water, the resulting

nanoparticles **NP1** were characterized by IR spectroscopy and solid-state ^{29}Si NMR to demonstrate successful functionalisation.

As shown in **Figure 4**, bands arising from the **Stalk** in the spectrum of **NP1** are observed at 1557 and 1498 cm^{-1} , which are not evident in the corresponding **NP** spectrum. In the region from 2700 to 3100 cm^{-1} , the spectrum of **NP1** is dominated by bands associated with CTAB at 2854 and 2924 cm^{-1} , and hence the spectral features of **Stalk** are not clearly observed, except for a shoulder at 2972 cm^{-1} . CTAB, initially retained within the pores to maximize functionalisation on the external surface, was then extracted with HCl to afford **NP2**. The molecular recognition sites on the **Stalk**, which were protonated during acidic extraction, were deprotonated (and hence re-activated) by suspending **NP2** in an aqueous solution of Et_3N . The mixture was stirred for 48 h at 70°C, before centrifugation and washing to give **NP3**. CTAB elimination was confirmed by the disappearance of its associated bands at 2854 and 2924 cm^{-1} in the IR spectrum of **NP3**.

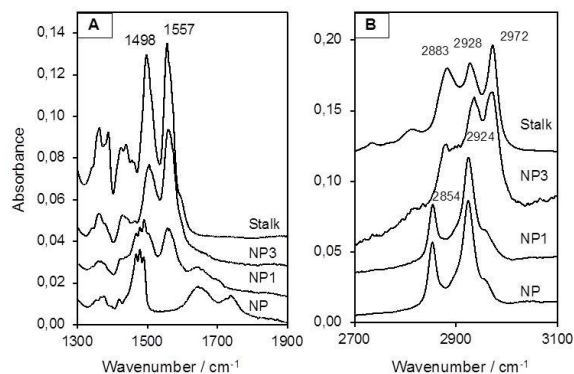


Figure 4. IR spectra (A) from 1300 to 1900 cm^{-1} and (B) from 2700 to 3100 cm^{-1} of NPs and the **Stalk** precursor

The solid-state ^{29}Si NMR spectrum of **NP1** (**Figure 5 (B)**) also demonstrated the successful grafting of the **Stalk**, with two sets of chemical shifts being evident. The first set, at -47, -57 and -67 ppm, are attributed to T^1 , T^2 and T^3 sites, respectively, arising from the **Stalk** and are not observed in the spectrum of **NP** (**Figure 5 (A)**), as expected. The relatively high intensity of the T^2 signal compared to that of the T^3 signal indicates that condensation of the ethoxy/hydroxy species on the **Stalk** is incomplete after grafting, with some of the four Si sites in the grafted **Stalk** moieties being bound to the surface of the nanoparticles *via* two oxo bonds instead of three. The second set of peaks in the ^{29}Si -NMR spectra, at -95, -100 and -109 ppm, are assigned to Q^2 , Q^3 and Q^4 species, respectively, within the MCM-41 framework. The increase in the relative intensity of the Q^4 signal for **NP1**, and the corresponding decrease in the $-\text{OH}:\text{Si}$ ratio for the Q^n species following grafting (from 0.75 to 0.65), is associated with conversion of Q^2 and Q^3 species to Q^4 species during functionalisation. Thermogravimetric analysis (TGA) confirmed the presence of the organic **Stalk** on the surface of the MCM-41 nanoparticles (**Figure S3**).

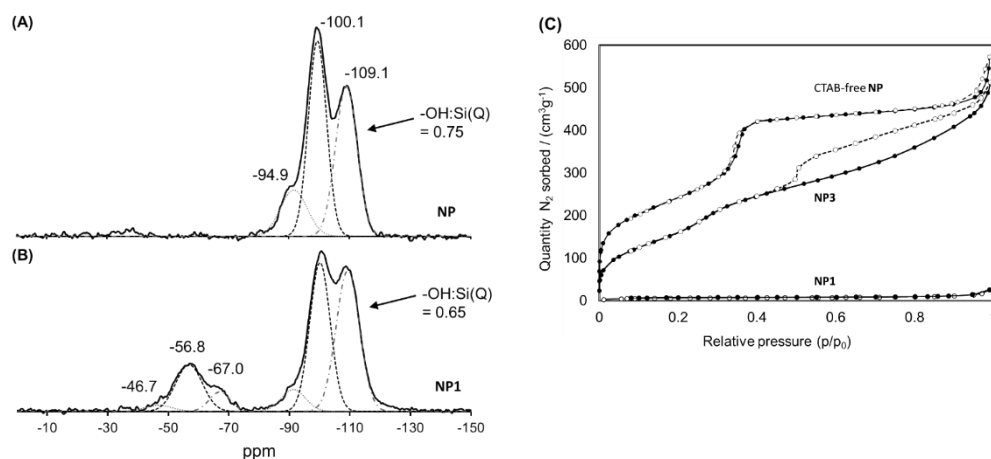


Figure 5. Solid state ^{29}Si NMR spectra of **NP** (A) and **NP1** (B). (C) Isotherms of CTAB-free **NP**, **NP1** and **NP3** obtained by nitrogen sorption analyses.

The N₂ adsorption/desorption isotherms of **NP1** (grafted nanoparticles containing CTAB) are compared with those of CTAB-free **NP** and **NP3** in **Figure 5 (C)**. As expected, **NP1** exhibits very low accessibility to small molecules such as N₂, with a corresponding BJH adsorption pore volume and BET surface area of 0.03 cm³.g⁻¹ and 21 m².g⁻¹, respectively, consistent with the presence of CTAB within the mesopores. In contrast, removal of CTAB by washing yields a mesoporous material (Type IV isotherm) with a pore volume of 0.92 cm³.g⁻¹ and BET surface area of 902 m².g⁻¹. The isotherm of CTAB-free NP shows a steep increase near P/P₀ = 0.35, consistent with the presence of ordered mesopores. The corresponding data for **NP3** are also consistent with a highly porous material, albeit with a lower porosity (0.72 cm³.g⁻¹) and surface area (675 m².g⁻¹) than the unfunctionalised material **NP**. A slight increase in the adsorption branch at p/p₀ around 0.3 for **NP3** is consistent with smaller pore size than in the CTAB-free **NP**. More significantly, the isotherm of **NP3** has a distinct hysteresis with non-parallel branches on the adsorption and desorption arms. Such profiles are often observed for porous samples having ink-bottle type pores, consistent with modification of the external surface in the vicinity of the pore openings. These data suggest that functionalisation leads to some obstruction of the nanoparticle porosity, although the pore network is still highly accessible to small molecules such as N₂.

To load the functionalised nanoparticles with a drug cargo, **NP3** was suspended with several different cargos, including RhB, PI or CPT in water or DMSO. **Cap** was then added to the suspension to complex the **Stalks** on the surface and thus block the pores. The suspension was stirred at RT for a further 48 h and then centrifuged. The resulting nanoparticles were vigorously washed with DMSO (until a transparent supernatant was obtained) to remove adsorbed cargo that had not been incorporated within the pores prior to capping and residual physisorbed **Cap**. After additional washings with water and EtOH, the nanoparticles were dried under vacuum to give the **NP3RhB**, **NP3PI** or **NP3CPT** nanomaterials, depending on the cargo compound used.

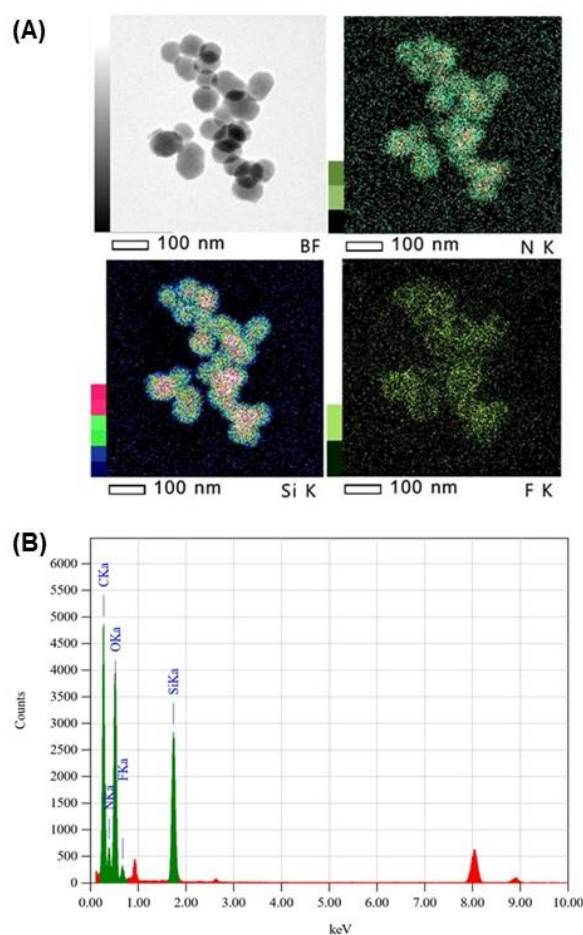


Figure 6. (A) Bright field (BF) and elemental mapping (Si, N and F) and **(B)** EDX spectrum of **NP3CPT** by EDX-TEM.

The quantity of **Cap** complexed by **Stalk** in the **NP3** system was assessed by monitoring the relative quantities of Si, N and F in the **NP3CPT** system via energy dispersive X-ray (EDX)-TEM analysis. The results obtained are illustrated in **Figures 6** and **S4** (ESI). The semi-quantitative EDX analysis revealed significant quantities of F atoms (**Figure S4**) arising from the presence of surface-complexed **Cap**. In addition, the relative quantities of Si, N and F atoms (**Figure S4**), together with the relative masses of SiO₂ and **Stalk** determined by TGA (**Figure S3**), are consistent with a

Stalk:Cap molar ratio of 3:1 and a CPT loading of 4.0 wt % in the **NP3CPT** system. The data also reveal the presence of three **Stalk** (and hence one **Cap**) per 100 Si atoms (see ESI). This is consistent with the expected complexation between the complementary **Stalk** and **Cap** and the associated obstruction of the MSNP pore apertures in **NP3CPT**. In addition, the size and morphology of the nanoparticles were characterized at different stages during the synthesis to verify their uniformity. SEM and TEM micrographs of **NP**, **NP3** and **NP3CPT** (**Figure 7A**) showed relatively mono-dispersed nanoparticles with diameters ranging from 110 to 200 nm. The TEM images also revealed the expected transverse porosity in the materials. A comparison of the nanoparticle size obtained by SEM with that observed in solution *via* DLS studies (**Figure 7B**) indicates that the functionalised nanoparticles are essentially unaggregated after being dispersed in solution, with relatively small polydispersity indices. As expected, the data indicate that functionalisation of **NP** with **Stalk**, to form **NP3**, results in a small increase in the hydrodynamic diameter of the nanoparticles (from 140 to ~190 nm) without any increase in polydispersity, consistent with functionalisation of the particles by **Stalk**. The size and low polydispersity of **NP3** remain unchanged after impregnation with CPT (to form **NP3CPT**).

Consequently, the resulting drug loaded **NP3CPT** have a suitable size for nanomedicine applications.

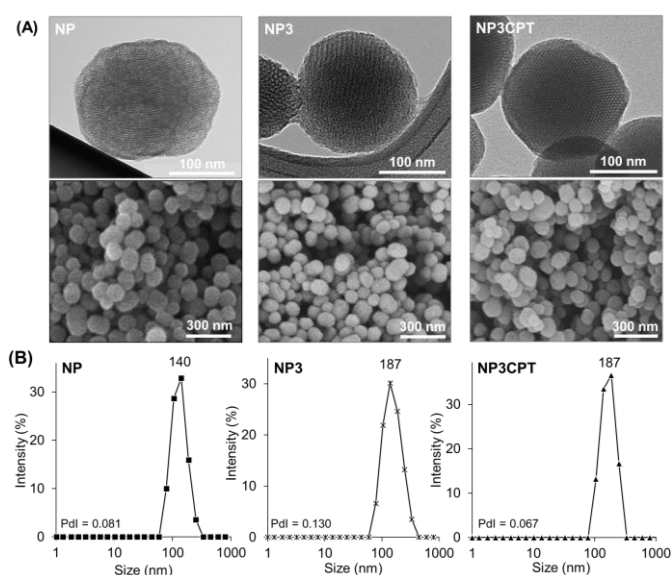


Figure 7. (A) TEM and SEM images and (B) DLS measurements of **NP**, **NP3** and **NP3CPT**, nanomaterials. The polydispersity indices (PDI, measured by DLS) are included.

Thereafter, the stability of the **Stalk-Cap** complex as a function of pH (2, 5.5 or 7.4) was investigated in the case of **NP3CPT** by monitoring CPT release in solution *via* UV-Vis absorbance at 380 nm (**Figure 8**). The data indicated that only about 5% of the drug is released at neutral pH, confirming that sufficient **Stalk-Cap** bonds remain intact to ensure that the nanoparticles can efficiently transport the chemotherapeutic agents within the blood stream without premature release of the payload.

In contrast, around 38 % of CPT was released at pH 5.5 after 24 h. This pH was chosen to mimic that present within the endosomal compartments, and the results suggest that release of the cargo and **Cap** would be expected to occur within the endosomal regions. As expected, essentially quantitative release of **Cap** is observed at pH 2 after only a few hours. This result demonstrates that the removal of the chemotherapeutic **Cap** and subsequent release of the second drug loaded within the pores is controlled by environmental pH. Our earlier studies of the release of **Cap** confirm that its release concentration profile is similar to that of CPT.

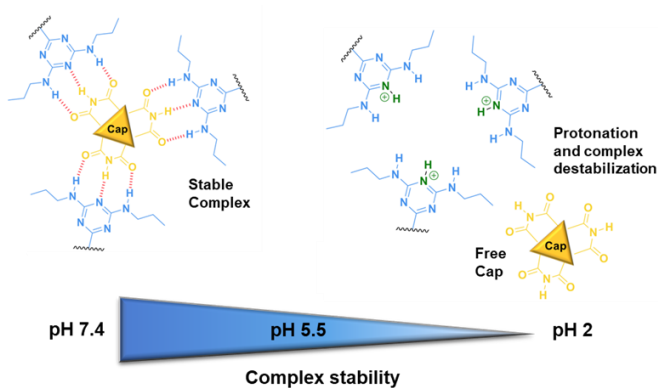
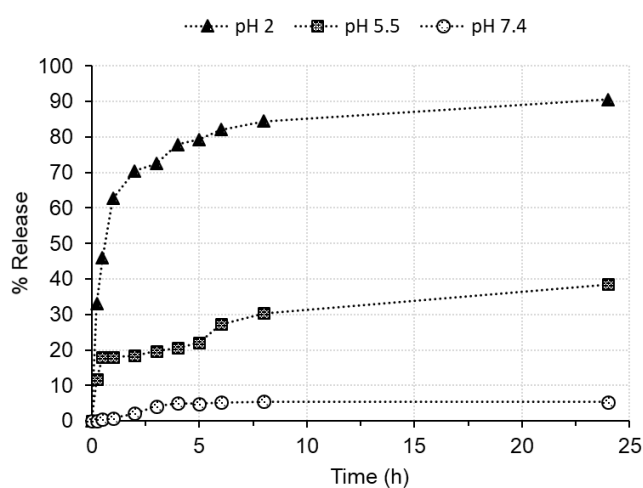


Figure 8. Release of CPT from NP3CPT mesopores (determined by UV/Vis absorbance spectroscopy) as function of pH

In vitro studies on MCF-7 breast cancer cells

Kinetic study

For kinetic studies, MCF-7 cells were incubated at 37 °C for varying time intervals with 40 $\mu\text{g}\cdot\text{mL}^{-1}$ of NP3PI. The nanoparticle internalization was detected by flow cytometry after co-staining with DAPI to exclude dead cells, and the results presented below represent the averages calculated from three independent experiments.

As shown in **Figure 9 (A)**, the internalization was relatively fast, with 50% of MCF-7 cells containing nanoparticles after 20 h. This result confirmed that the size and nature of the nanoparticles were suitable for penetrating MCF-7 cancer cells. In addition, it is important to note that, for this measurement, only living cells were considered, and since our cap is biologically active, it might lead to the death of some cells that would be not detected by the approach used here. Hence, the values reported in **Figure 9 (A)** represent a lower limit for the apparent rate of internalisation.

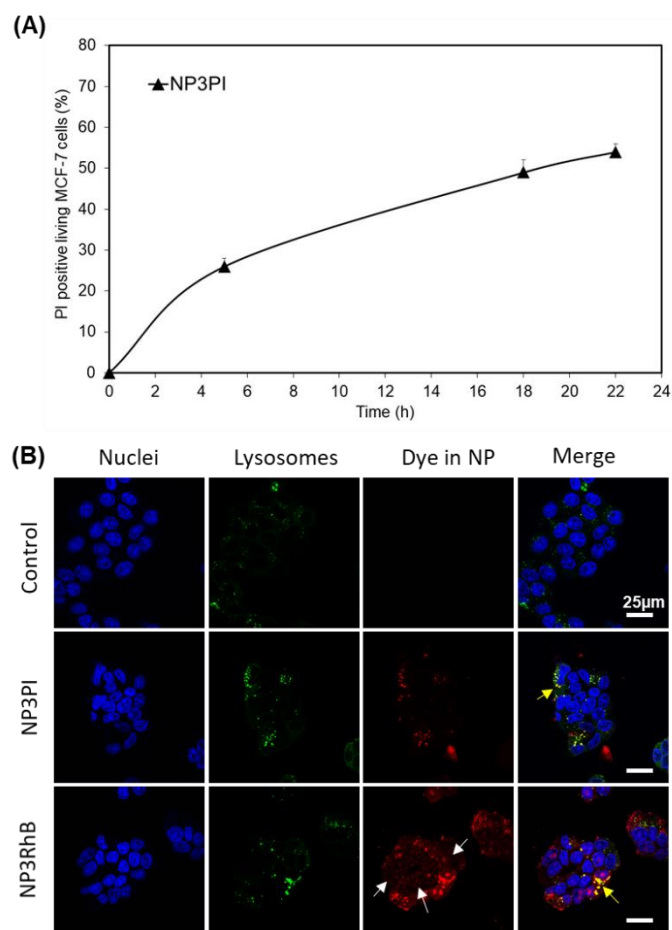


Figure 9. (A) Internalization rate in MCF-7 cells using NP3PI. Error bars show standard deviation. (B) Confocal microscopy of Control (top), NP3PI (middle) and NP3RhB (bottom) after 20 h of incubation in a culture medium of MCF-7 cells

Uptake of nanoparticles in MCF-7 cells - Confocal Microscopy

To assess the uptake of nanoparticles in MCF-7 breast cancer cells, the cells were incubated for 20 h with $40 \mu\text{g}\cdot\text{mL}^{-1}$ of NP3RhB and NP3PI (Figure 9 (B)). The endo-lysosomal compartments were labelled with the green lysotracker and the nuclei marked in blue with the Hoechst stain. The Control images correspond to cells that have not been incubated with nanoparticles. In culture cells incubated with NP3PI, we can detect the presence of PI in red, and the yellow regions observed in the merged picture demonstrate the co-localization of NP3PI with endo-lysosomal compartments (see yellow arrows in Figure 9 (B)). This validates the internalisation of nanoparticles by the endocytosis pathway. In addition, when culture cells were incubated with nanoparticles loaded with RhB (NP3RhB), which is known to cross the cell membranes (and lysosome), we observed that the red dye fully stained the cytoplasm (see white arrows in Figure 9 (B)), confirming the release of the cargo molecules once the nanocarriers have entered the lysosomes. Overall, these data confirm (1) the uptake of the nanomaterials by the cell *via* the endo-lysosomal pathway and (2) that the release of active drug molecules is triggered at lysosomal pH, consistent with the pH-sensitivity data included in Figure 8.

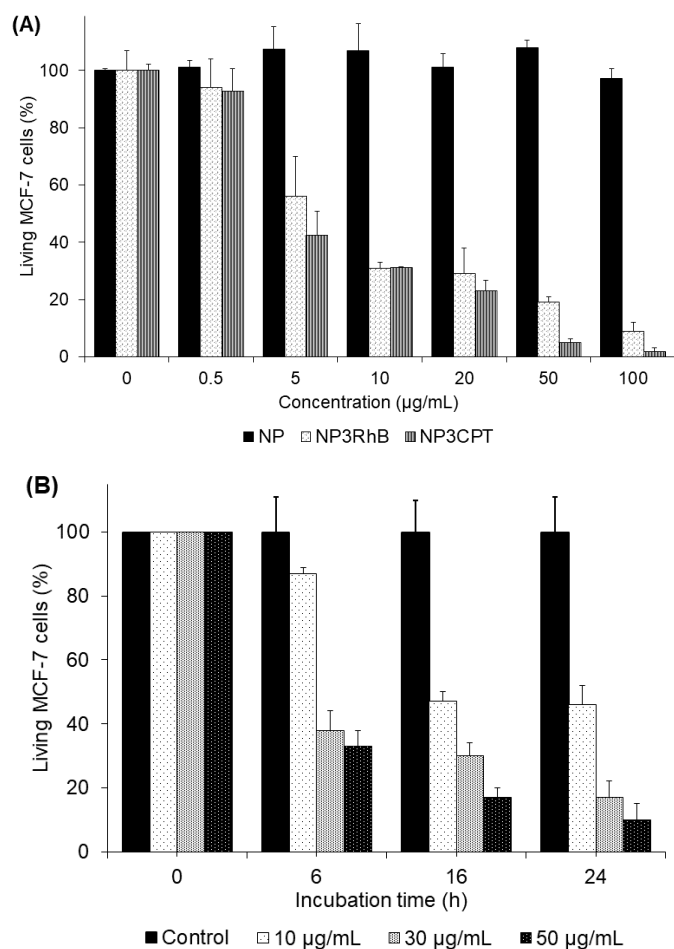


Figure 10. (A) Cytotoxicity test of NP, NP3RhB and NP3CPT on MCF-7 cells after 72 h of incubation. (B) Cytotoxic effect of NP3CPT on MCF-7 as a function of incubation time and concentration. Error bars show standard deviation.

Cytotoxicity

The cytotoxicity of the new nanocarrier on MCF-7 breast cancer cells was then assessed *via* MTT assays. This study was performed with NP3RhB to determine the action of the cytotoxic 5-FU cap (RhB is known to be inactive²⁹) and with NP3CPT to determine if the cytotoxicity is increased by the presence of two different drugs. As shown in **Figure 10 (A)**, which illustrates cell death after 72 h of incubation with different batches of nanoparticles, NP exhibited essentially no cytotoxic activity (Control). The corresponding data for NP3RhB confirmed that the active cap was released, with good cytotoxicity demonstrated against MCF-7 cells. Finally, NP3CPT showed excellent tumour elimination, with more than 90% of cell death from a concentration of 50 µg.mL⁻¹ and up to 99% of cell death at 100 µg.mL⁻¹. The IC₅₀ of NP3CPT under these conditions is 4 µg.mL⁻¹, a particularly low value, which demonstrates the complementary and concomitant activity of the multidrug approach employed here, where each drug attacks the cells *via* different and complementary biological pathways.

Following this promising result, the activity of NP3CPT on MCF-7 cells was determined at shorter times. For this, incubations with NP3CPT at 10, 30 and 50 µg.mL⁻¹ were performed and followed by MTT assays after 6, 16 and 24 h. The results presented in **Figure 10 (B)** revealed that the action of NP3CPT was relatively fast. Indeed, after only 6 h of incubation, MTT assay results already showed a significant inhibition of cell growth. Moreover, after 24 h the results obtained were close to those observed on the previous study carried out after 72 h of incubation. These data confirm that essentially complete apoptosis of the MCF-7 cells can be achieved, *in vitro*, with the NP3CPT system within 24 h.

Conclusions

Herein, a silica-based nanocarrier system in which CPT was loaded within the pores and a 5-FU analogue complexed on the surface as a pore-capping agent has been shown to be a viable system for environmentally-triggered,

sequential release of these anti-cancer drugs for successful eradication of breast cancer cells. The nanosystem has been shown to respond to changes in environmental pH, thus enabling the delivery of a multi-drug payload within the lysosomal compartment of cancer cells. In contrast, almost no release of the cytotoxic compounds was observed under normal physiological conditions, thus minimising toxic side-effects in normal tissue and within the blood stream.

The efficacy of this approach was demonstrated through *in vitro* trials, involving the consecutive delivery of the 5-FU derivative (rapid release) and CPT (diffusion-limited release) in MCF-7 breast cancer cells.

Under the conditions used in this study, essentially complete breast cancer cell apoptosis was observed within 72 h. This result highlights the attractive synergistic effect of the two drugs acting in tandem against tumoral cells. Additionally, the loading of the two synergistic drugs into silica nanocarriers offers the significant advantage of preventing the debilitating side effect usually associated with combination cancer therapy.

It is envisaged that the approach developed in this work could also be extended to multi-combinatorial therapy, involving the sequential release of three or more drugs.

Acknowledgements

We are grateful to the CNRS and the French Higher Education and Research Ministry. This work was partially funded with a young researcher grant from the French National Research agency (ANR-2010-JCJC-1006-01). We acknowledge the imaging facility MRI, member of the national infrastructure France-BioImaging supported by the French National Research Agency (ANR-10-INBS-04, «Investments for the future»). We thank Prof. Fuyuhiko Tamanoi for access to EDX-HRTEM at Kyoto University and help and support in pH-release testing.

Notes and references

1. Q. Hu, W. Sun, C. Wang and Z. Gu, *Advanced drug delivery reviews*, 2016, **98**, 19-34.
2. R. Bayat Mokhtari, T. S. Homayouni, N. Baluch, E. Morgatskaya, S. Kumar, B. Das and H. Yeger, *Oncotarget*, 2017, **8**, 38022-38043.
3. P. Sharma and James P. Allison, *Cell*, 2015, **161**, 205-214.
4. O. C. Farokhzad and R. Langer, *Advanced Drug Delivery Reviews*, 2006, **58**, 1456-1459.
5. N. Kolishetti, S. Dhar, P. M. Valencia, L. Q. Lin, R. Karnik, S. J. Lippard, R. Langer and O. C. Farokhzad, *Proceedings of the National Academy of Sciences*, 2010, **107**, 17939-17944.
6. A. Jhaveri, P. Deshpande and V. Torchilin, *Journal of Controlled Release*, 2014, **190**, 352-370.
7. J. K. Patra, G. Das, L. F. Fraceto, E. V. R. Campos, M. d. P. Rodriguez-Torres, L. S. Acosta-Torres, L. A. Diaz-Torres, R. Grillo, M. K. Swamy, S. Sharma, S. Habtemariam and H.-S. Shin, *Journal of Nanobiotechnology*, 2018, **16**, 71.
8. R. K. Singh, J. C. Knowles and H.-W. Kim, *Journal of Tissue Engineering*, 2019, **10**, 2041731419877528.
9. I. I. Slowing, B. G. Trewyn, S. Giri and V. S.-Y. Lin, *Advanced Functional Materials*, 2007, **17**, 1225-1236.
10. I. I. Slowing, J. L. Vivero-Escoto, C.-W. Wu and V. S. Y. Lin, *Advanced Drug Delivery Reviews*, 2008, **60**, 1278-1288.
11. J. M. Rosenholm, V. Mamaeva, C. Sahlgren and M. Lindén, *Nanomedicine*, 2012, **7**, 111-120.
12. Y. Wang, Q. Zhao, N. Han, L. Bai, J. Li, J. Liu, E. Che, L. Hu, Q. Zhang, T. Jiang and S. Wang, *Nanomedicine: Nanotechnology, Biology and Medicine*, 2015, **11**, 313-327.
13. J. Wen, K. Yang, F. Liu, H. Li, Y. Xu and S. Sun, *Chemical Society Reviews*, 2017, **46**, 6024-6045.
14. M. Vallet-Regí, F. Balas and D. Arcos, *Angewandte Chemie International Edition*, 2007, **46**, 7548-7558.
15. C.-H. Lee, S.-H. Cheng, I.-P. Huang, J. S. Souris, C.-S. Yang, C.-Y. Mou and L.-W. Lo, *Angewandte Chemie International Edition*, 2010, **49**, 8214-8219.
16. L. Yuan, Q. Tang, D. Yang, J. Z. Zhang, F. Zhang and J. Hu, *The Journal of Physical Chemistry C*, 2011, **115**, 9926-9932.
17. C. Théron, A. Gallud, S. Giret, M. Maynadier, D. Grégoire, P. Puche, E. Jacquet, G. Pop, O. Sgarbura, V. Bellet, U. Hibner, J. I. Zink, M. Garcia, M. Wong Chi Man, C. Carcel and M. Gary-Bobo, *RSC Advances*, 2015, **5**, 64932-64936.
18. Q. Zhang, F. Liu, K. T. Nguyen, X. Ma, X. Wang, B. Xing and Y. Zhao, *Advanced Functional Materials*, 2012, **22**, 5144-5156.
19. A. Rahikkala, S. A. P. Pereira, P. Figueiredo, M. L. C. Passos, A. R. T. S. Araújo, M. L. M. F. S. Saraiva and H. A. Santos, *Advanced Biosystems*, 2018, **2**, 1800020.
20. Z.-Y. Li, Y. Liu, X.-Q. Wang, L.-H. Liu, J.-J. Hu, G.-F. Luo, W.-H. Chen, L. Rong and X.-Z. Zhang, *ACS Applied Materials & Interfaces*, 2013, **5**, 7995-8001.
21. F. Muhammad, M. Guo, A. Wang, J. Zhao, W. Qi, Y. Guo and G. Zhu, *Journal of Colloid and Interface Science*, 2014, **434**, 1-8.
22. M. C. Llinàs, G. Martínez-Edo, A. Cascante, I. Porcar, S. Borrós and D. Sánchez-García, *Drug Deliv*, 2018, **25**, 1137-1146.
23. S. Giret, C. Théron, A. Gallud, M. Maynadier, M. Gary-Bobo, M. Garcia, M. Wong Chi Man and C. Carcel, *Chemistry – A European Journal*, 2013, **19**, 12806-12814.

24. L. F. Liu, S. D. Desai, T.-K. Li, Y. Mao, M. E. I. Sun and S.-P. Sim, *Annals of the New York Academy of Sciences*, 2000, **922**, 1-10.
25. D. B. Longley, D. P. Harkin and P. G. Johnston, *Nature Reviews Cancer*, 2003, **3**, 330-338.
26. S. Guichard, I. Hennebelle, R. Bugat and P. Canal, *Biochem Pharmacol*, 1998, **55**, 667-676.
27. Q. Luo, P. Wang, Y. Miao, H. He and X. Tang, *Carbohydrate Polymers*, 2012, **87**, 2642-2647.
28. C. Théron, A. Birault, M. Bernhardt, L. M. A. Ali, C. Nguyen, M. Gary-Bobo, J. R. Bartlett, M. Wong Chi Man and C. Carcel, *Journal of Sol-Gel Science and Technology*, 2019, **89**, 45-55.
29. P. Fisher, *Wildlife Society Bulletin (1973-2006)*, 1999, **27**, 318-329.