Size controlled protein nanoemulsions for active targeting of folate receptor positive cells

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A B S T R A C T
Bovine serum albumin (BSA) nanoemulsions were produced by high pressure homogenization with a tri-block copolymer (Poloxamer 407), which presents a central hydrophobic chain of polyoxypropylene (PPO) and two identical lateral hydrophilic chains of polyethylene glycol (PEG). We observed a linear correlation between tri-block copolymer concentration and size – the use of 5 mg/mL of Poloxamer 407 yields nanoemulsions smaller than 100 nm. Molecular dynamics and fluorescent tagging of the tri-block copolymer highlight their mechanistic role on the size of emulsions. This novel method enables the fabrication of highly stable albumin emulsions in the nano-size range, highly desirable for controlled drug delivery. Folic Acid (FA)-tagged protein nanoemulsions were shown to promote specific folate receptor (FR)-mediated targeting in FR positive cells. The novel strategy presented here enables the construction of size controlled, functionalized protein-based nanoemulsions with excellent characteristics for active targeting in cancer therapy.

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1. Introduction

Particle size is a feature of particular importance for intravenous therapeutic applications. Nanoparticles intended for systemic circulation should be large enough to prevent their rapid leakage into blood capillaries but small enough to escape capture by macrophages in the reticuloendothelial system, such as in the liver and spleen [1,2]. Nanoparticles for drug delivery smaller than 100 nm can be intravenously administered and are taken up more efficiently by cells [3].

Albumin is widely used to prepare protein nanoparticles, which can be easily functionalized chemically [4]. Functionalized nanoparticles can be utilized for targeted drug delivery, improving cellular uptake and ameliorating undesired toxic side effects associated with conventional chemotherapeutic regimens [3,5].

The state of the art on the production and mechanism of formation of protein-based capsules was recently reviewed by our group [6]. The production of albumin-based nanoparticles can be achieved using specialized nanotechnology techniques like desolvation, emulsification, thermal gelation, and more recently through nano-spray drying, nab-technology and self-assembly [4,6–8]. Susslick and Grinstaff, were pioneers in developing a methodology using high-intensity ultrasound, which is a high-energy emulsification method, to produce aqueous suspensions of proteinaceous micro-capsules [9]. It was demonstrated that high shear forces applied to biphasic water/hydrophobic solvent systems can induce conformation changes in proteins which tend to migrate to the water/solvent.
interface, stabilizing the microcapsules [6,10]. In a previous study, our group demonstrated that the use of a large globular protein, BSA, with clear distinct hydrophilic and hydrophobic areas, n-dodecane (as an organic solvent) and a ratio of 95% of aqueous phase/5% of n-dodecane subjected to high-intensity ultrasound, resulted in small and stable microspheres. However, the particles produced using this method never presented size smaller than 250 nm [10].

Here, we describe a novel method based on high pressure homogenization (also a high-energy emulsification method) rather than using high-intensity ultrasound, of a biphasic system (aqueous phase and vegetable oil) to produce stable protein nanoparticles of small, defined and controllable size. Taking into account the production method, where an aqueous solution and oil are emulsified by high pressure homogenization, we denominated these nanoparticles as nanoemulsions. The addition of a PEGylated surfactant (Poloxamer 407) regulates the particle size while the chemical functionalization of albumin with FA confers specificity toward FR positive cells, thus achieving active targeting.

2. Material and methods

2.1. Nanoemulsion preparation

The preparation of nanoemulsions was achieved using a high pressure homogenizer (APV-2000, Denmark). BSA (Sigma–Aldrich, USA) previously dissolved in phosphate buffered saline (PBS, salt concentration 150 mM) was emulsified with an organic solvent (vegetable oil) by subjecting the mixture to varying number of homogenization cycles at high pressure. Different concentrations of BSA (1: 5; 10 and 50 mg/mL), of Poloxamer 407 (Sigma–Aldrich, USA) and different ratios of protein solution/vegetable oil (95:5; 99:5/0.5) were tested in order to determine how these factors influence the properties of the resulting particles. PBS concentration was maintained constant throughout the study to avoid the influence of salt concentration in the production optimization.

A fluorescent agent was introduced in the aqueous phase by using Fluorescein isothiocyanate (FITC) labelled-BSA (Sigma–Aldrich, USA) at an initial formulation of a 1:20 ratio (m/m). Previously prepared BSA-FA conjugate solution was added at 1:100 ratio (m/m) to the aqueous phase to obtain nanoemulsions with FA (the targeting agent) on their surface.

2.2. Conjugation of FA with BSA

In brief, FA (Sigma–Aldrich, USA) was dissolved in PBS, pH 7.4, and ‘activated’ with 5-fold excess of N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (Sigma–Aldrich, USA) and a 1-fold excess of N-hydroxysulfosuccinimide (Sigma–Aldrich, USA) for 2 h at room temperature. To this activated FA was then added 10 mg/mL of BSA and left to react for 8–10 h, using a 75:1 FA to BSA moles ratio. The excess FA and other reactants were removed from the conjugated protein by dialyzing against PBS for 8 days. After dialysis, quantification of FA linked to BSA was determined by spectroscopic measurements at the maximum wavelength of 350 nm. Using a calibration curve of different concentrations of FA, the concentration of FA linked to BSA was determined. The quantitation of BSA was performed with Bio-Rad protein assay (Bio-Rad laboratories, GmbH) based on the shift of absorbance maximum for an acidic solution of Coomassie Brilliant Blue G-250 at 595 nm when binding to protein occurred. Therefore, it was found that 1.625 mg/mL of FA was linked to 7 mg/mL of BSA, which corresponds to 35 moles of FA per 1 mole of BSA.

2.3. Determination of size, zeta-potential and long-time stability

The nanoemulsions were analyzed for their size distribution and their zeta-potential (ζ-potential), at pH 7.4 (PBS buffer) and at 25 °C, using dynamic light scattering in a Malvern zetasizer NS, by photon correlation spectroscopy (PCS) and by electrophoretic laser Doppler anemometry, respectively. The protein concentration was kept constant at 10 mg/mL. The values for viscosity and refractive index were taken as 0.890 cP and 1.330, respectively. Each sample was measured in triplicate and results are presented as mean value ± standard deviation.

Nanoemulsions, in suspension, were stored at 4 °C for a period of 20 weeks. After predetermined storage times, the stability parameters, particles size, polydispersity and zeta-potential, were determined as described above.

The nanoemulsion’s stability was also evaluated in biological medium through the incubation in PBS buffer containing 10% of fetal bovine serum (FBS) at 37 °C for 24 h.

2.4. Determination of nanoemulsion formation efficiency

The amount of BSA used in nanoemulsion production was quantified by measuring the amount of free protein with the Bradford protein assay (Bio-Rad Laboratories, USA), using BSA solutions as standards. After synthesis, the nanoparticles-containing phase was collected by centrifugation (1000 × g for 45 min) using centricon tubes (Amicon Ultra-15, Millipore), a centrifugal filter unit containing a cellulose membrane with a molecular weight cut-off of 100 KDa. The free protein in the aqueous phase after separation was quantified and the efficiency of nanoemulsion formation was determined using the formula:

\[
\text{Nanoemulsion formation (\%) = } \frac{[C_i] - [C_f]}{[C_i]} \times 100
\]

where \([C_i]\) and \([C_f]\) is the initial and final concentration of the protein in the aqueous solution, respectively. Each sample was assayed in triplicate.

2.5. Determination of entrapment efficiency of different pharmacological drugs

The hydrophilic drugs were dissolved in aqueous phase and the hydrophobic oil phase of the initial formulation, which was posteriorly subjected to high pressure homogenization. The initial concentration of each drug was: 5-Fluorouracil = 788 μg/mL; doxorubicin = 50 μg/mL; methotrexate = 688.5 μg/mL; taxol = 2.5 μg/mL; carbon monoxide releasing molecule-2 (CORM-2) = 100 μg/mL. Calibration curves with different concentrations of each hydrophilic drug were performed by spectroscopic measurements at the maximum wavelengths (5-Fluorouracil: 265.5 nm; doxorubicin: 481 nm; methotrexate: 302 nm) and the respective LOD were calculated to be: 5-Fluorouracil = 0.0446 μg/mL; doxorubicin = 0.0362 μg/mL; methotrexate = 0.0443 μg/mL according to [11]. After nanoemulsion production, a separation of phases was performed by centrifugation (1000 × g for 45 min) using centricon tubes (Amicon Ultra-15, Millipore). The free drug in the aqueous phase was quantified and the entrapment efficiency was determined using the formula:

\[
\text{Entrapment efficiency (\%) = } \frac{[\text{drug}]_{\text{Total}} - [\text{drug}]_{\text{Free}}}{[\text{drug}]_{\text{Total}}} \times 100
\]

where \([\text{drug}]_{\text{Total}}\) and \([\text{drug}]_{\text{Free}}\) is the total concentration of drug added in formulation and the free concentration of drug in aqueous phase solution, respectively.
2.6. Morphology of nanoemulsions

The morphology of the BSA nanoemulsions was examined by fluorescence microscopy and transmission electron microscopy (TEM) imaging systems. The nanoemulsion solution was analyzed in a Leica DM5000B fluorescence microscope in order to determine if fluorescently labeled protein was present at the nanoparticle’s surface. For TEM analyses, nanoemulsion samples were applied to glow discharged carbon-coated copper grids followed by negative staining with a solution of 1% (w/v) Uranyl acetate and then analyzed in a JEOL JEM2100F transmission electron microscope.

2.7. Molecular dynamics simulation

The simulations were performed with the software package GROMACS [12], using Martini force-field [13–15]. The system size was chosen according to the minimum image convention taking into account a cut-off of 1.2 nm. The bonds lengths were constrained with LINCS [16]. Non-bonded interactions were calculated using a twin-range method, with short and long range cut-offs of 0.9–1.2 nm, respectively. Neighbor searching was carried out up to 1.2 nm and updated every ten steps. A time step of integration of 5 fs was used. A reaction field correction for the electrostatic interactions was applied using a dielectric constant of 15. Pressure control was implemented using the Berendsen barostat [17], with a reference pressure of 1 bar, 3.0 ps of relaxation time and isothermal compressibility of $3.0 \times 10^{-5}$ bar$^{-1}$. Temperature control was set using the Berendsen thermostat [17] at 300 K. Each component of the system was included in separate heated bath, with temperatures controlling constants of 0.30 ps. Two replica simulations of 250 ns in length were carried out using different initial velocities taken from a Maxwell–Boltzman distribution at 300 K. Initially the components were distributed uniformly through the simulation box and allowed to interact freely during the simulation. The simulations were extended to 250 ns and not more because the system had reached an equilibrium. No significant changes were observed, in both the energy and position and distribution of the particles in the system, in the last nanoseconds of the simulations.

2.8. Particle preparation, using ultrasound methodology, for confocal observations

In order to produce larger sized particles to facilitate confocal analysis, the bottom of a high-intensity ultrasonic horn was positioned at the interface of the protein aqueous solution and vegetal oil and activated at an amplitude of 40% at 10°C (±1) for 3 min. The ratio of aqueous/organic phase (%) used were 90/10, and 5 mg/mL of BSA and 5 mg/mL of Poloxamer 407 were introduced in the aqueous phase. BSA-FITC and Poloxamer 407-rhodamine B solution, previously prepared, were used for particle preparation.

2.9. Conjugation of rhodamine B with poloxamer 407

Poloxamer 407 coupled with rhodamine B (Sigma–Aldrich, USA) was synthesized using the Steglich esterification method [18]. Briefly, rhodamine B (0.2 mM) preactivated with N,N-dicyclohexylcarbodiimide (DCC) (Sigma–Aldrich, USA) (0.2 mM) was conjugated to the hydroxyl (OH) groups of Poloxamer (0.1 mM) in dichloromethane (DCM) (20 mL) including 4-(dimethylamino) pyridine (DMAP) (0.2 mM), triethylamine (0.1 mL), and pyridine (0.1 mL) (Sigma–Aldrich, USA). The solution was stirred at room temperature for 1 day. After the reaction, the solution was filtered and the solvent evaporated under reduced pressure. The resulting light pink solid was then dissolved in PBS and the excess rhodamine B and other reactants were removed from conjugated poloxamer using benzoylated dialysis tubing (Sigma–Aldrich, MWCO 2 KDa). Dialysis was performed against PBS, pH 7.4, for 3 days. The dialysis was stopped when no rhodamine B was detected in the dialysis buffer.

2.10. Cells and culture conditions

Human skin fibroblasts (BJ5ta cell line) (ATCC, CRL-4001) and a subclone of HeLa cells [19] (KB cell line) (ATCC, CCL-17) were obtained from American Type Culture Collection (LGC Standards, UK). Exponentially growing cultures were maintained in a humidified atmosphere of 5% CO₂ in air at 37 °C. All culture media and supplements were purchased from Sigma–Aldrich. Adherent cell lines (BJ5ta and KB cells) were grown in T75 flasks (TPP, Switzerland).

<table>
<thead>
<tr>
<th>Cell line</th>
<th>Media</th>
<th>Supplements</th>
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<tbody>
<tr>
<td>BJ5ta</td>
<td>4 Parts of Dulbecco’s modified eagle’s medium (DMEM)</td>
<td>2 mM l-glutamine; 4.5 mg/mL glucose; 1.5 mg/mL sodium bicarbonate; 10% (v/v) of FBS; 1% (v/v) of penicillin/streptomycin solution;</td>
</tr>
<tr>
<td></td>
<td>1 Part of M199 Medium</td>
<td>10 μg/mL of hygromycin B</td>
</tr>
<tr>
<td>KB</td>
<td>RPMI-1640 Medium</td>
<td>10% (v/v) FBS; 1% (v/v) penicillin/streptomycin solution; 2 mM l-glutamine; 2 mg/mL sodium bicarbonate</td>
</tr>
</tbody>
</table>

2.11. Cell viability assay

Cell viability was studied using the Promega CellTiter 96® AQueous non-radioactive cell proliferation (MTS) assay (Promega, USA). BJ5ta cells were seeded in 96-well tissue culture polystyrene plates (TPP, Switzerland) at a density of $1.1 \times 10^4$ cells/well and incubated overnight to promote cell adhesion. The cells were incubated with different concentrations of nanoemulsions (from 75 up to 900 μg/mL) and following incubation for 24, 48 and 72 h, a MTS mixture was then added and the cells were further incubated for 4 h at 37 °C. After this period, the plates were placed on Spectra MAX 340PC microwell plate reader and the absorbance of the formazan product was read at 940 nm. Cell viability was expressed as a percentage relative to the negative control (untreated control cells).

2.12. Quantification of oxidative stress -2′-7′-dichlorofluorescein assay

BJ5ta cells were seeded in 12-well plates (TPP, Switzerland) at a density of $1 \times 10^5$ cells/well and, after overnight cellular adhesion, incubated with different concentrations of nanoemulsions (from 75 up to 900 μg/mL) for 12, 24, and 48 h. After each period of incubation, cells were washed twice with PBS buffer and incubated with 200 μL of a 100 μM DCFH₂-DA (Molecular Probes by Life Technologies, USA) solution for 30 min in the dark at 37 °C. Extracellular DCFH₂-DA was removed by washing the cells twice with PBS buffer. Hydrogen peroxide ($H₂O₂$) was then added to the positive control in this phase (1:5 (v/v)). Lysis solution (90% DMSO solution) 10% PBS buffer (v/v) was added and cells incubated on a shaker for 10 min in a dark at room temperature. The lysates were transferred in to black opaque 96-well plates and fluorescence intensity was then read using wavelengths of 485 nm excitation and 525 nm emission, at 25 °C, using a Fluoroskan Ascent FL plate reader.

2.13. Confocal microscopy

BSA nanoemulsions containing FITC labeled-BSA and FA–BSA conjugate at their particle’s surface were prepared and their inter-
nalization by cells was analyzed after separation of unbound FITC–BSA and FA–BSA using centrifugation. The FR positive cancer cell line KB cell line was used in this study. KB cells were seeded at a density of $5 \times 10^4$ cells/well and incubated overnight to promote cell adhesion. Solutions of nanoemulsions with and without FA fusions were prepared in FA-free Hank's Balanced Salt solution (HBSS medium) (Lonza, Belgium). Cells were washed twice with PBS buffer and then 300 µg/mL nanoemulsion suspensions were added to the monolayer, which was then incubated at 37°C for 2 h. After three further washing steps with cold PBS buffer, cells were fixed with 500 µL of paraformaldehyde 4% (v/v) for 30 min and permeabilized with Triton 0.1% (v/v) solution. After two washing steps with PBS, the fixed cells were incubated with 5 units/mL of Alexa Fluor 568 Phalloidin (Molecular Probes by Life Technologies, USA). The coverslips were placed on slides coated with 2 µL of Permafluor (Thermo Scientific, United Kingdom) containing 5 µg/mL of Hoechst (Molecular Probes by Life Technologies, USA).

Microscopic observations were performed using an inverted Zeiss confocal laser scanning microscope (CLSM; Olympus Fluoview FV1000).

BSA emulsions containing FITC labeled-BSA and rhodamine B labeled-Poloxamer 407 and prepared using the ultrasound method were also observed by confocal laser scanning microscopy.

Three dimensional models were produced with the software Imaris image analysis program.

2.14. Cell uptake quantification

KB cells were seeded in 6-well plates (TPP, Switzerland) at a density of $2 \times 10^4$ cells/well and incubated overnight to promote cell adhesion. Solutions of nanoemulsions with and without FA fusions were prepared in HBSS medium. Cells were washed twice with PBS pH 7.4 and nanoemulsion suspensions at a concentration of 300 µg/mL were added to the monolayer of cells and incubated at 37°C for 2 h. After three further washing steps with cold PBS buffer, cells were lysed with one solution of Tris–HCl buffer at 10 mM and the fluorescence of these solutions was then analyzed with an Horiba Jobin Yvon Spectrofluorog-3 spectrofluorimeter using the $\lambda_{exc} = 480$ nm, scanning within a wavelength range of 490–650 nm. Calibration curves of each solution of tested nanoemulsions were also measured.

3. Results and discussion

3.1. High pressure homogenization methodology for producing stable and size controlled nanoemulsions

For the production of functionalized nanoemulsions of a defined size, we used a high pressure homogenizer. High pressure systems are continuously operated and droplet disruption is achieved through a combination of turbulent flow and laminar flow producing shear stresses and cavitation [20]. Different emulsification steps are usually involved in preparing nanoparticles by high pressure homogenizer and, in some cases, the addition of cross-linking agents is necessary [21–25]. The strategy described here involves a single step of emulsification which allows high batch-to-batch reproducibility, ease of scale-up, simplicity, resulting in homogeneously small, highly stable protein nanoemulsions. In a previous study, we demonstrated that the hydrodynamic cavitation phenomenon involved in high pressure homogenizer creates negligible amounts of radicals, when compared with ultrasound [26]. The applied pressures and the number of homogenization cycles can affect particle size and stability in the final emulsion. Stable emulsions were obtained when using a pressure of 600 bar, in the first stage of pressure, and 250 bar, in the second stage of pressure and, when applying 26 homogenization cycles. The ratio between...
Table 1

<table>
<thead>
<tr>
<th>Concentration range with viability higher than 90% (µg/mL)</th>
<th>Oxidative stress levels in exposed cells (DCF assay)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nanoemulsions (ratio 95/5)</td>
<td>n.d.</td>
</tr>
<tr>
<td>Nanoemulsions (ratio 99.5/0.5)</td>
<td>Similar to control cells</td>
</tr>
<tr>
<td>Nanoemulsions (ratio 99.5/0.5) + Poloxamer 407</td>
<td>Similar to control cells</td>
</tr>
</tbody>
</table>

*Nanoemulsions were prepared using BSA solution at 10 mg/mL and oil with specified ratios and with 5 mg/mL of Poloxamer 407 in aqueous solution when indicated.

BSA solution and organic solvent may also influence the size of the resulting nanoemulsion particles, as observed when using ultrasonic procedures [10]. Two different BSA/vegetable oil ratios, 95%/5% and 99.5%/0.5%, were tested using different BSA concentrations (1; 5; 10 and 50 mg/mL). The size and stability of these samples over time were analyzed (Fig. 1A). Nanoemulsions prepared using high concentrations of protein (10 and 50 mg/mL) were formed with high efficiency (above 90%) and displayed smaller sizes and higher stability over time (Fig. 1A). Nanoemulsions prepared with lower vegetable oil content exhibited constant size (Fig. 1A), polydispersity and zeta-potential values (data not shown) for 20 weeks.

Cytotoxicity evaluation using immortalized human normal skin fibroblasts further supported that 99.5% BSA solution at 10 mg/mL/0.5% vegetable oil was ideal for nanoemulsion preparation. While some cytotoxicity was induced in fibroblasts exposed to nanoemulsions prepared with higher oil content, no cytotoxicity whatever was detected in cells contacting for at least 72 h with nanoemulsions prepared with a low oil content (Table 1). From this point onwards, the nanoemulsions were always prepared using 99.5% BSA solution at 10 mg/mL/0.5% vegetable oil.

3.2. Poloxamer 407 concentrations have a direct effect on nanoemulsion size

The inclusion of hydrophobic non-ionic surfactant (Poloxamer 407) was investigated as a means to obtain homogeneously small nanoemulsions with stealthy features, required for systemic administration. Poloxamer 407 is a tri-block copolymer with a central hydrophobic chain of polyoxypropylene (PPO) and two identical lateral hydrophilic chains of polyethylene glycol (PEG) (Fig. 1B). A linear correlation was found between surfactant concentration and nanoemulsion particle size (Fig. 1C), illustrated by dynamic light scattering. TEM imaging confirmed these results and revealed spherical shaped BSA based nanoemulsions (Fig. 1D). These results are in agreement with a previous characterization using STEM imaging, reported by us [27].

The physicochemical characterization of nanoemulsions containing Poloxamer 407 showed that, in addition to the precise control of nanoemulsion particle size, the presence of the PEGylated surfactant also determines their surface characteristics. The introduction of 5 mg/mL of Poloxamer 407 into the formulation resulted in nanoemulsion particles smaller than 100 nm and with a zeta-potential close to zero. Even after 20 weeks of storage in PBS at 4 °C, the properties of these formulations remained unaltered (Fig. 2), demonstrating their high stability as previously described [27].

We have tested the stability of nanoemulsions containing Poloxamer 407 when exposed to biological medium (see Section 2 for details). Minor variations (<10%) in the size of the nanoemulsions were observed after 24 h of incubation while the respective zeta-potential values remained unaltered. Given the high ionic strength of the biological medium and the high salt and content, these results demonstrated the nanoemulsions’ high stability [28].

Poloxamers are known to bind to the surface of the nanospheres through hydrophobic interaction of the PPO chains, whilst the hydrophilic PEG chains extend into the surrounding medium and create a steric barrier to form a protective coating [29–33]. We suggest that the mechanism of nanoemulsion formation, as well as its tunable properties such as size, arise from the chemical nature of both the Poloxamer and the BSA protein, and the subsequent physicochemical interactions between them. Thus, the steric barrier resulting from hydrophilic PEG chains of Poloxamer may prevent the interaction between an approaching nanoparticle and the target cell. PEG molecules can also reduce the adsorption of opsonins and other serum proteins through a mechanism known as the steric repulsion effect [32,34]. PEG molecules will form a dynamic molecular “cloud” over the particle surface, due to the chain flexibility and the electrical neutrality of the PEG backbone [32,34,35]. A correlation between surface charge and opsonization has also been demonstrated in vitro, indicating that neutral charged particles have a much lower clearance from circulation than electrically charged particles [36]. In a previous study, we demonstrated that nanoparticles containing 2.5 mg/mL of PEGylated surfactant are more readily recognized and internalized by macrophages than nanoparticles containing 5 mg/mL of surfactant [27]. These results showed that the presence of PEGylated surfactant at a concentration of 5 mg/mL appears to promote PEG chains to sufficiently extend into the aqueous solution to shield the surface of the nanoparticles and improve resistance to macrophage uptake. Therefore, this neutral and hydrophilic surface in PEGylated nanoemulsions can generate stealth particles that are less phagocytized, resulting in a longer half-life in systemic circulation [27].

Molecular dynamics simulation was used to better understand the interactions between Poloxamer 407 and BSA with the oil phase, as experimentally, it is very difficult to clarify the interaction between components. The results obtained clearly illustrate the interactions between the components of the system (Fig. 3A). The hydrophobic part of Poloxamer 407 (blue = polyoxypropylene part) interacts preferably with the oil, decreasing hydrophobic interactions between the oil and the protein, which may be responsible for nanoemulsion size reduction upon surfactant addition.

Upon analyzing the radial distribution of the components (Fig. 3B), an additional tool to clarify the relative positions of the components, we may speculate that the polyoxypropylene component is in close proximity to the oil whilst the polyoxyethylene interacts preferentially with the protein. The overall tri-block surfactant seems to make a matrix with the protein surrounding the
oil, which is in line with the results obtained by the zeta-potential measurements that indicated that nanoemulsions without Poloxamer 407 exhibit negatively charged particles while the presence of Poloxamer 407 leads to a neutral particle surface.

An emulsion using BSA labeled with FITC and Poloxamer 407 labeled with rhodamine B was prepared using ultrasounds, which also involves high shear forces, so as to obtain larger particles suitable for confocal microscopy. 3D Reconstruction of the images obtained with the confocal microscope confirmed that the nanoemulsion particle shells were composed of protein and PEGylated surfactant (Fig. 3C). The PEGylated surfactant is found preferentially at the oil interface but also on the surface of the protein shell, with both components interacting in order to form the shell of the nanoemulsion particles, as suggested by the zeta-potential measurements.

3.3. Biological characterization of nanoemulsion formulations

In order to evaluate the cytotoxicity of nanoemulsion formulations containing Poloxamer 407, cell viability was assessed in human skin fibroblasts (BJ5ta cell line) using the MTS assay. Nanoemulsions with surfactant did not induce toxicity at the wide range of concentrations tested, over a period of 72 h of contact (Table 1). Cell viability was also evaluated at 24 and 48 h (data not shown). One important potential risk of using nanostructured formulations is the possibility of eliciting high levels of intracellular reactive oxygen species (ROS), causing the deleterious oxidation of biomolecules. To evaluate if the nanoemulsion formulations induce the production of intracellular ROS, the 2′,7′-dichlorofluorescein (DCF) assay was used. In this assay, intracellular ROS oxidate the non-fluorescent DCFH$_2$ to a highly fluorescent compound, DCF, which is then quantified. The intensity of the fluorescent signal detected by this method is proportional to the amount of ROS produced within the cells. Nanoemulsion formulations did not induce oxidative stress in human BJ5ta cells (Table 1).

3.4. Production and characterization of FA-PEGylated BSA nanoemulsions

For evaluating FR-mediated cell targeting, nanoemulsions containing imaging and targeting agents were prepared by introducing BSA–FITC and BSA–FA conjugates, respectively. BSA–FITC was introduced into the initial formulation, containing 5 mg/mL of Poloxamer 407, at a 1:20 ratio (m/m) and BSA–FA at a 1:100 ratio (m/m) relative to the BSA protein. Fluorescence microscopy observations and quantification of FA, using RIDASCREEN®FAST Folic acid test kit, demonstrated that fluorescently labeled BSA and a high FA concentration were present at the nanoemulsion’s surface (633.2 ± 114.39 µg/L). The physicochemical characterization of nanoemulsions containing FA was performed and we observed that the values of size, PDI and zeta-potential are similar to that of non-targeted PEGylated nanoemulsions.

Subsequent studies, including cellular internalization of tagged nanoemulsions, were performed using these PEGylated nanoemulsions containing 5 mg/mL of Poloxamer 407. Their small size (under 100 nm) and stealth characteristic make them ideal nanoparticles for intravenous application. The higher concentration of Poloxamer 407 induce a better stealth, which might reduce the interaction...
between nanoparticle and the cell, decreasing their unspecific cellular uptake [38,39]. This phenomenon favors specific cell uptake mediated by folate.

Specific cellular internalization of functionalized nanoemulsions was then evaluated in FR positive KB cell line [40,41]. FA conjugates typically exhibit high binding affinities for FR (KD = 1–10 nM), which is abundantly overexpressed in many malignant tumors – for example, up to 80 pmol (mg membrane protein) in KB cells. The quantification of internalization by spectrofluorimetry demonstrated that the FA-PEGylated nanoemulsions are internalized by cells more efficiently and to a higher extent (5 fold) than non-targeted PEGylated nanoemulsions (Fig. 4A).

3D reconstructions of images obtained by confocal microscopy also showed the specific internalization of FA-nanoemulsions with the involvement of the cell membrane (Fig. 4B). Similar specificity in FA-mediated cell uptake of these nanoemulsions was verified in other FR-alpha positive cell lines, MDA-MB 468 [42,43] and Caco-2 [44,45] (data not shown). While for these cancer cell lines no competitive studies were done, in primary activated human macrophages, which are known to express FR-beta, these were performed using an anti-human FR-beta antibody. When the FR-beta was preliminarily blocked with the FR antibody, no significant differences in cell internalization were observed between FA-PEGylated nanoemulsions and non-targeted PEGylated nanoemulsions. However, experiments with the unblocked FR demonstrated a 40% increase of the internalization for the FA-targeted nanoemulsions. The comparison between FA-targeted nanoemulsions internalization with and without FR blocking also demonstrated differences, with a significant inhibition of internalization observed when the FR was blocked.

These results suggest that FA-PEGylated nanoemulsions interact efficiently with FR at the cellular surface and promote FR-mediated endocytosis.

3.5. Drug delivery applications

As this versatile system is intended for drug delivery purposes, we performed the entrapment of different pharmacological drugs, which were successfully incorporated in the nanoemulsions (Table 2).

The general characteristics of the nanoemulsions were not significantly altered by drug loading, although the entrapment efficiency was different according to the drug included. Entrapment efficiency of lipophilic drugs may be considered to be 100% as the oil phase is totally incorporated in the nanoemulsions. The percentage of entrapped drug was smaller in the case of hydrophilic drugs. Additionally, we performed a study of drug release for nanoemulsions loaded with methotrexate. 20 days after nanoemulsion preparation, a phase separation was performed to separate the free drug from the encapsulated fraction using PD-10 desalting columns (GE Healthcare). The quantification of methotrexate in the aqueous phase separated demonstrated that approximately 43% of the drug had been released. All these results of the entrapment of drugs and the structural studies of these nanoemulsions suggest that they can efficiently incorporate hydrophobic compounds.
Table 2
Characterization of nanoemulsions loaded with different pharmacological drugs. Nanoemulsions were prepared using 99.5% aqueous solution containing BSA at 10 mg/mL and Poloxamer 407 at 5 mg/mL / 0.5% vegetable oil. (n.d. = not determined).

<table>
<thead>
<tr>
<th>Compound structure</th>
<th>Compound solubility</th>
<th>Z-average [d, nm] (PDI)</th>
<th>Entrapment efficiency (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nanoemulsions with 5-Fluorouracil</td>
<td>Hydrophilic</td>
<td>n.d.</td>
<td>26.0 ± 3.5</td>
</tr>
<tr>
<td>Nanoemulsions with Doxorubicin</td>
<td>Hydrophilic</td>
<td>84.57 ± 0.73 (0.130 ± 0.004)</td>
<td>47.3 ± 6.4</td>
</tr>
<tr>
<td>Nanoemulsions with Methotrexate</td>
<td>Hydrophilic</td>
<td>84.51 ± 0.36 (0.166 ± 0.002)</td>
<td>25.0 ± 2.6</td>
</tr>
<tr>
<td>Nanoemulsions with Taxol</td>
<td>Lipophilic</td>
<td>93.85 ± 0.71 (0.136 ± 0.017)</td>
<td>n.d.</td>
</tr>
<tr>
<td>Nanoemulsions with CORM-2</td>
<td>Lipophilic</td>
<td>89.86 ± 4.24 (0.143 ± 0.016)</td>
<td>n.d.</td>
</tr>
</tbody>
</table>

* Entrapment efficiency of lipophilic drugs may be considered to be 100% as the oil phase is totally incorporated in the nanoemulsions.

These novel FA-PEGylated BSA nanoemulsions thus arise as promising vehicles for selective delivery of drugs to a target cell population that overexpresses FR.

4. Conclusions

In conclusion, we have developed a high pressure homogenization procedure that enables the formation of size controlled, stable and functionalized BSA nanoemulsions. The amount of protein and the hydrophobic phase volume were demonstrated to be crucial parameters in the nanoemulsion formation process, as previously described for ultrasound method [10]. By combining this novel production method with the introduction of Poloxamer 407, protein-based nanoemulsions with controllable size were developed. The inclusion of FA in the formulation allowed the functionalization of such vectors for FA-mediated active targeting of FR positive cells. Their small size, high stability over time and absence of cytotoxicity, together with their specific targeting capacity in cells expressing FR, are ideal characteristics to support their application in drug delivery into diseased tissues. The procedure here described can be used to produce virtually any type of functionalized protein-based nanoemulsions increasing their scope as a vehicle for targeted drug delivery purposes.

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