Complete Disaggregation of MCF-7-derived Breast Tumour Spheroids with Very Low Concentrations of α-Mangostin Loaded in CD44 Thioaptamer-tagged Nanoparticles

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Abstract

Background: α-Mangostin (αMG) is a natural substance that exerts a wide range of antitumor effects. Recently, we described that free αMG was able to dissociate multicellular tumour spheroids (MCTSs) generated from breast carcinoma cells and to reduce their cellular viability and motility. Here, αMG was encapsulated into lipidic nanoparticles (NPs), conjugated or not to a CD44 thioaptamer, and the anticancer action evaluated against MCF-7 breast MCTSs.

Methods: NPs containing αMG were formulated with a core of polylactic-co-glycolyc acid. Some of them were decorated with a CD44 thioaptamer using as catalysts 1-ethyl-3- (3-dimethylaminopropyl) carbodiimide and N-hydroxysuccinimide. Both size and density of MCF-7-derived MCTSs were monitored during 72 h of treatment with NPs carrying 0.1, 0.5 and 1.0 µg/ml final concentrations of αMG. MCTSs were cultured on Matrigel or gelatine to better simulate the extracellular environment.

Results: The NPs without thioaptamer and conveying 0.1 µg/ml αMG caused a significant dissociation of the MCTSs grown in gelatine after 24 h of treatment (p < 0.01). The most significant disaggregation of MCTSs was obtained using NPs carrying 0.5 µg/ml αMG (p < 0.01). A similar dissociating effect was observed when MCTSs were cultured in Matrigel under the same conditions for 48 – 72 h. By contrast, only concentrations over 1.0 µg/ml of free αMG were able to provoke a damage to MCTSs, consisting in a substantial reduction in their size (p < 0.05). Since the MCTS dissociation induced by αMG-loaded NPs occurred only in the presence of Matrigel or gelatine, an impairment of cell contacts to collagen fibres was likely responsible of this effect. Finally, the treatment of MCTSs with αMG-loaded NPs that were conjugated to the CD44 thioaptamer caused a similar decrease in density but a lower expansion of the spheroid, suggesting that a significant number of cells were died or arrested in cycle.

Conclusion: Very low concentrations of αMG delivered by lipidic NPs are sufficient to provoke a substantial disaggregation of MCF-7 MCTSs that involves cell-to-collagen contacts. Similarly, the treatment of MCTSs with NPs conjugated to a CD44 thioaptamer leads to MCTS dissociation but through a more damaging action that causes also a reduction in cell number.

Key words: α-mangostin, multicellular tumour spheroid, breast cancer cell line, MCF-7, nanoparticle, thioaptamer.

Introduction

α-Mangostin (αMG), a xanthone that is contained in large amount in Garcinia mangostana Linn, has been demonstrated to possess several antitumor properties under in vitro and in vivo conditions [1]. The wide range of pharmacological activities of αMG and the low frequency of its adverse
and covered by lipophilic/polyethylene glycol layers of nanoparticles (NPs) as a vehicle for intravenous strategies that have been investigating is the use of pharmacodynamics of antitumor drugs are fields on "enhanced permeability and retention" (EPR) effect, the body is increased [11]. In addition, according to the and hence the lifespan of the transported drug in the are not recognized by the reticular endothelial system as a suitable tool to reduce the dosage of the drug NPs targeting specific tumour cells can be considered as a suitable tool to reduce the dosage of the drug cargo and, therefore, the occurrence of adverse effects of chemotherapy.

According to these concepts, we generated MCTSs by a MCF-7 breast carcinoma cell line and the antitumor effects of αMG, as a free compound or encapsulated in lipidic NPs, were evaluated. We found that very low concentrations of αMG delivered by NPs caused a significant reduction in spheroid compactness, without increasing cell invasiveness. Since MCF-7 cells largely express CD44, the antitumor activity of αMG-loaded NPs conjugated to a CD44 thioaptamer was also investigated.

Materials and Methods

Materials

Reagents were purchased from Sigma-Aldrich (St. Louis, MO, USA), unless otherwise stated.

MCF-7 cell monolayer

MCF-7 human breast carcinoma cell line (the European Collection of Authenticated Cell Cultures, ECAAC) were expanded in culture flasks under standard conditions (37 °C, 5% CO2, 95% humidity) or subjected to a low oxygen tension (1% O2, 94% N2, 5% CO2). Cells were cultured in complete Dulbecco’s modified Eagle medium (DMEM), containing 2% fetal calf serum (v/v), 100 U/ml penicillin and 100 µg/ml streptomycin. The culture medium was routinely replaced twice a week. For cell expansion, subconfluent cells were detached using a solution of 0.05% trypsin in 0.53 mM EDTA. MCF-7 cells were cultured also as monolayers on 96-well flat-bottomed plates at the density of 1x10^4 cells/well.

Generation and morphological analysis of MCTSs

For the initial characterization of the spheroids, MCF-7 cells were seeded in ultra-low attachment (ULA) 6-well flat-bottomed plates (Corning; Sigma-Aldrich) at densities ranging from 1x10^3 to 2x10^4 cells/ml. The culture medium was serum-free DMEM/F12 (1:1 v/v) containing 2% B27 (ThermoFisher Scientific, Waltham, MA, USA), 20 ng/ml epidermal growth factor (EGF, Peprotech, Rocky Hill, NJ, USA), 20 ng/ml basic fibroblast growth factor (bFGF, Peprotech), 100 U/ml penicillin, and 100 µg/ml streptomycin (MCTS medium). To perform a high throughput screening, 200 µl of cell suspension in MCTS medium was seeded on each well of ULA 96-well round-bottomed plates (Corning) and to encourage cell aggregation, the plates were centrifuged at 100 g for 3 min. Under this condition, spheroids of diameter ranging between 250 and 300 µm were generated after three days. Then, some
MCTSs were treated with different concentrations of αMG dissolved in 0.1% (v/v) dimethylsulfoxide (DMSO), while control MCTSs were exposed only to the vehicle.

Images were captured using an inverted microscope (IX50, Olympus Italia, Segrate, Italy), equipped with an Olympus camera, and imported into Image-J software (Fiji, http://imagej.nih.gov). Using the graphic utilities of Image-J, the largest (equatorial) border of each spheroid was manually drawn and the pseudo-circular area of its phase-contrast image was calculated. After background subtraction, the MCTS density was measured using the “mean grey” command of the software applied to the 8-bit inverted images of each spheroid. This parameter provided the grey half-tone intensity-to-area ratio of the pseudo-circular images of MCTSs. Morphological measurements were performed each day of treatment and expressed as percentage of those obtained at day 3 after cell seeding (100%). Some spheroids grew for three days on ULA plates and then were transferred to adherent 96-well flat-bottomed plates (Corning) that were coated with 100 µl of 0.1% (w/v) gelatine or Matrigel (Corning) for better simulating the extracellular matrix (ECM) environment of solid tumours [17].

Cell viability assay

MCF-7 cells at the density of 1x10^4 cells/well were seeded on adherent 96-well flat-bottomed plates and cultured as monolayers in complete medium for 48 h [18]. Then, PrestoBlue Cell Viability Reagent 10X solution (ThermoFisher Scientific) was added in each well at a 10% final concentration. Fluorescence was measured at 530 nm excitation and 590 nm emission on a Wallac VICTOR² multiwell plate reader (Perkin Elmer, Milan, Italy) 4 h after dye addition.

Production of NPs

Lipid-polymer combinational nanoparticles were synthesized from polylactic-co-glycolic acid (PLGA), soybean lecithin and 1,2-diasteroyl-glycero-3-phosphoethanolamine-N-carboxy(polyethylene glycol)2000 (DSPE-PEG2000–COOH) using a nanoprecipitation technique combined with self-assembly [19]. PLGA (50:50, M.W. 30-70 kDa) was first dissolved in acetonitrile at a concentration of 1 mg/ml. Lecithin and DSPE-PEG2000–COOH were dissolved with a 7:3 molar ratio in 4% ethanol aqueous solution at 15% of the PLGA polymer weight and heated to 65 °C. Then, the PLGA/acetonitrile solution was added to the lipid/aqueous solution in drop-wise manner, followed by vortexing for 3 min. αMG was added to the PLGA solution in the ratio 1:10 (w/w) relative to the polymer. The solution was then subjected to indirect sonication for 5 min in ice-cold water. The NPs were allowed to self-assemble and the organic solvent to evaporate with continuous stirring for 3.5 h. The remaining organic solvent and the free molecules were removed by washing the NP solution two times in ethanol/water and the third time in distilled water using an Amicon Ultra-4 centrifugal filter with a molecular weight cut-off of 10 kDa (Millipore; Sigma-Aldrich). The NPs were sonicated again, filtered by 200 nm cut-off (Corning), and divided in two aliquots of 1 ml each. An aliquot was air dried for about 1 h, weighed, and dissolved in acetonitrile plus 30% phosphate buffer saline (PBS), pH 7.2, for the spectrophotometric measurement of drug encapsulation (peak of absorbance of αMG at 320 nm). The second aliquot of 1 ml was immediately used to perform the conjugation with the aptamer. The remained suspension of NPs was stored at -20 °C. Fluorescent nanoparticles were also prepared by adding 10 µl of the fluorochrome Nile Red at the concentration of 1 mg/ml.

Conjugation of the thioaptamer to NPs

A DNA thioaptamer that specifically binds to CD44 according to the formulation T6A6 described by Somasunderam et al. [20] was synthesized by Trilink Biotechnologies (San Diego, CA, USA). The 73-mer oligonucleotide is a back bone-modified aptamer in which the non-bridging phosphoryl oxygens are substituted with sulphur. This change of structure renders the aptamer more resistant to the cellular degradation exerted by nucleases. The thioaptamer (M.W. 24,229.0) was linked to an amino group at the 5' end to allow the carboxy group of DSPE-PEG2000-COOH to form a carbamide covalent bond. For this purpose, we used the two catalysts 1-ethyl-3- (3-dimethylaminopropyl) carbodiimide (EDC) and N-hydroxysuccinimide (NHS) according to the manufacturer’s instructions (ThermoFisher Scientific). Briefly, NPs (10 mg) were dissolved in 1 ml of distilled water, filtered with an Amicon Ultra-4 10 kDa (Millipore) and resuspended in 1 ml of buffered 0.1 M 2-(N-morpholino)ethanesulfonic acid, 0.5 M NaCl, pH 6.0, containing a molar concentration of EDC and NHS 10-fold and 25-fold higher than DSPE-PEG2000-COOH, respectively. NP suspension was shacked gently at room temperature for 15 min. The resulting NHS-activated NPs were washed twice in distilled water with Amicon Ultra-4 10 kDa filters, resuspended in 1 ml PBS and conjugated to 3 nmoles of thioaptamer corresponding to a 1:100 ratio with respect to 0.85 mg DSPE-PEG2000-COOH used for the synthesis of 10 mg NPs. To this purpose, 30 µl of 100 µM thioaptamer in PBS was added to 1 ml PBS containing 10 mg NHS-activated NPs and stirred at
room temperature for 2 h. To remove the free thioaptamer, the final solution was washed three times in distilled water, using the Amicon Ultra-4 10 kDa filter, and the NPs resuspended in distilled water at the concentration of 10 mg/ml.

**Characterization and size measurement of NPs**

NP size was measured by tuneable resistive pulse sensing (TRPS) using the qNano Nanoparticle Analyser V2.0 (iZON Science, Christchurch, New Zealand) equipped with a fluid cell module of 300 nm nanopores (NP300) and operated by the V3.1 Control Suite software.

The ability of NPs, free or conjugated to the thioaptamer, to enter cells was evaluated by phase contrast and fluorescence microscopy (Olympus IX50) using CD44 + human bone marrow mesenchymal stromal cells (BM-MSCs; TebuBio, Magenta, Italy) at the density of 4 x 10\(^3\)/ml. NPs were also observed by scanning electron microscopy (JSM-5200; JEOL, Tokyo, Japan).

**Western blotting analysis**

MCF-7 cells were homogenized in a glass tissue grinder in 20 mM HEPES, pH 7.5, containing 5 mM dithiothreitol, 2 mM EDTA, 0.1% CHAPS detergent, 0.1% Triton X-100, and protease inhibitors; then, they were centrifuged at 15,000 g for 15 min. The supernatant was diluted in loading buffer (2% SDS, 5% glycerol, 0.002% bromophenol blue, 4% β-mercaptoethanol in 0.25 M Tris-HCl, pH 6.8) and denatured by boiling for 3 min. Aliquots corresponding to 80 μg protein were analysed by SDS-PAGE (7.5% gel). Proteins were transferred onto a nitrocellulose membrane for 1 h. This membrane was then saturated with 5% dry milk for 1 h, washed with Tris-buffered saline, and probed overnight at 4°C with the specific primary antibodies (1:500 mouse monoclonal anti-HIF-1α and 1:1,000 mouse monoclonal anti-β-actin; Santa Cruz Biotechnology, Heidelberg, Germany). The membrane was then incubated for 1 h with the secondary antibody (1:2,500 horseradish peroxidase-conjugated species-specific anti-IgG; Santa Cruz Biotechnology).

**Statistical analysis**

Values are expressed as mean ± standard error of the mean (SEM). Linear regression analysis and one-way analysis of variance (ANOVA) followed by Bonferroni multiple-comparison test were performed using GraphPad Prism 4.0 software (San Diego, CA, USA). P < 0.05 was considered statistically significant.

**Results**

**Viability of MCF-7 cells treated with free αMG**

MCF-7 cells at the density of 1x10\(^4\) cells/well were cultured as monolayers on 96-well flat-bottomed plates under normoxic conditions. The treatment with 0.1-20 μg/ml αMG for 48 h caused a bi-modal and dose-dependent decrease in cell viability. Figure 1 shows that cells were markedly damaged by concentrations of αMG up to 1.0 μg/ml, while higher doses up to 20 μg/ml led to a less accentuated drop in viability.

![Figure 1. Effects of free αMG on the MCF-7 cell monolayer. The viability of 1x10\(^4\) MCF-7 cells was reduced after treatment with free αMG for 48 h in a dose-dependent manner. The cells were greatly affected by 0.1 - 1 μg/ml αMG that caused a sharp decrease in their viability. A further damage to cells was observed using higher concentrations of αMG, although their responsiveness was reduced. Values were subjected to linear regression analysis and are expressed as mean ± SEM of duplicated experiments.](image)

**Effects of free αMG against MCTSs**

As preliminary characterizing set of experiments, MCTSs were produced in ULA 6-well plates by seeding 1x, 5x, 10x and 20x10\(^3\) MCF-7 cells/ml and the increasing size of the spheroids with cell density was verified (Figure 2a). Then, to obtain a high-throughput availability of MCTSs, a single spheroid per well was grown in ULA round-bottomed 96-well plates for 15 days (Figure 2b). The seeding density of 1x10\(^4\) cells/well was chosen because represented the best condition to obtain spheroids that were not too large for the limited size of the well but enough compact and well-shaped to perform appropriate morphological investigations.
Figure 2. Effects of free αMG on MCTS size and density. (a) MCF-7 cells seeded at 1x, 5x, 10x and 20x10³ cells/ml in ULA 6-well flat-bottomed plates generated MCTSs of increasing size after 7 days. The spheroids merged into larger aggregates at the highest cell density (magnification: 50 X). (b) The images show a single MCTS per well that was grown in ULA 96-well round-bottomed plates for 15 days (representative images of triplicate experiments). The optimized seeding density of 1x10⁴ cells/well allowed to obtain 3D bodies with a circular profile and homogeneous cellular distribution (magnification: 50 X). The spheroids treated with free αMG for 24 h (c) and 48 h (d) were negatively affected by doses higher than 0.8-1.0 µg/ml (n = 2). MCTSs generated in the presence of 1.0-20 µg/ml free αMG showed a smaller area and a parallel increase in density with respect the untreated MCTSs. (e) Representative phase-contrast images of the MCTSs treated for 48 h with increasing doses of free αMG (magnification: 50 X). The numerical values correspond to the concentrations of αMG expressed as µg/ml. At the highest dose, a darker region of cellular necrosis is clearly visible. The treatment with free αMG produced damages to the spheroids that were shown as changes in equatorial area and density (Figure 2c, 2d). The lowest concentrations of αMG that significantly reduced the MCTS area were 3 µg/ml and 1 µg/ml, administered for 24 h (p < 0.05) and 48 h (p < 0.01), respectively. Conversely, the MCTS density underwent a parallel increase that reached statistical significance after treatment with αMG at the concentrations of 10 µg/ml for 24 h (p < 0.05) and 3 µg/ml for 48 h (p < 0.05). The irregular edges of the MCTSs and the homogenous distribution of their density observed at the highest doses (Figure 2e) were considered as further signs of αMG toxicity.

Characterization and cell uptake of NPs

The mean diameter of NPs was 227.0 ± 88 nm and 174.0 ± 29 nm before and after filtration, respectively (Figure 3a, 3b). Both diameter and size distribution of NPs were consistent with the images obtained by SEM (Figure 3c). The fluorescent dye Nile Red was loaded in NPs to trace their uptake inside cells. BM-MSCs were chosen to better perform this test because they are larger than MCF-7 after adhesion on plates of polystyrene. BM-MSCs were incubated with NPs and observed by a combination of phase contrast and fluorescence microscopy. The process of NP incorporation into cells was effective after 24 h (Figure 3d, 3e) and larger aggregates of NPs were generated after 48 h of incubation (Figure 3f).
Figure 3. Characterization and cell uptake of NPs. (a) The size distribution of NPs measured just after their production was 150-380 nm. (b) The range of NP diameter was restricted to 110-260 nm after filtration, with a mean value of 174.0 ± 29 nm. (c) NP size and distribution were consistent with the images obtained by SEM. (d, e) BM-MSCs were incubated with Nile Red-loaded NPs and contemporary observed by phase contrast and fluorescence microscopy after 24 h. (f) Most NPs fused into larger aggregates after 48 h. Images are representative of duplicate experiments.

Figure 4. Morphological changes of MCTSs grown in Matrigel and treated with NPs. NPs loaded with αMG were able to dissociate the spheroids. MCTS treatment for 48-72 h with NPs that were not conjugated to the thioaptamer (NP[-]) and carrying 0.1 or 0.5 μg/ml αMG led to cell detachment after 48-72 h. NPs conjugated to the thioaptamer (NP[+]) and carrying 0.1 μg/ml αMG reduced the size of the spheroids, while NPs loaded with 0.5 μg/ml αMG provoked MCTS disaggregation. The images are representative of two separate experiments (magnification: 50X).

The amount of αMG loaded in NPs was 1.4 μg/mg. A suspension up to 1 mg/ml of NPs without αMG in complete medium did not affect growth and viability of cells, irrespective of the presence or absence of the thioaptamer (data not shown).
Disaggregation of MSCTs cultured in Matrigel and treated with αMG-loaded NPs

MCTSs were grown in Matrigel and their morphology was observed after 24, 48 and 72 h of treatment with NPs (Figure 4). The concentration of 0.1 µg/ml αMG carried by NPs without thioaptamer (NP[-]) was sufficient to loosen MCTS cells in a time-dependent manner. The dose of 0.5 µg/ml αMG also provoked a robust dissociation of MCTSs, independently of the absence or presence of the thioaptamer. The administration of NPs conjugated to the thioaptamer (NP[+]) at 0.1 µg/ml αMG reduced the size of MCTSs and produced a lower effect on cell aggregation.

Disaggregation of MSCTs cultured in gelatine and treated with αMG-loaded NPs

αMG conveyed by NPs caused the disaggregation of MCTSs also when they were grown in gelatine (Figure 5). An initial drop in spheroid density was observed just after 24 h of treatment with 0.1 µg/ml αMG. Maximal spheroid dissociation was obtained using NPs with 0.5 µg/ml αMG for 72 h, as it was shown by the lowest density and the highest area of the MCTSs. The presence of the thioaptamer linked to NPs (NP[+]) attenuated the MCTS expansion after treatment with 1.0 µg/ml αMG for 48 h (p<0.01) and 72 h (p<0.05), as it was shown by the lower values of the area relative to NP[-]. However, the reduction in density produced by NP[+] was of the same magnitude of that obtained with NP[-] at the corresponding αMG concentrations.

Figure 5. Morphological changes of MCTSs grown in gelatine and treated with NPs. The treatment with αMG-loaded NPs increased the area of MCTSs (upper slopes) and caused a parallel drop in their density (lower slopes). The lowest dose of αMG that provoked a decrease in MCTS density was 0.1 µg/ml just after 24 h of treatment. The expansion of the area was less pronounced using NPs conjugated to the thioaptamer (NP[+]) than that caused by NPs without thioaptamer (NP[-]). *p < 0.05 and **p < 0.01, comparing the same concentrations of αMG carried by NP[+] vs NP[-]. *p < 0.05 and **p < 0.01, comparing NP[-] vs the corresponding untreated MCTSs. *p < 0.05 and **p < 0.01, comparing NP[+] vs the corresponding untreated MCTSs. Micrographs show the dramatic changes of MCTS morphology after treatment with NPs at concentrations of αMG up to 1.0 µg/ml delivered for 72 h (magnification: 50X). The numerical values under the images are concentrations of αMG expressed as µg/ml.
Reduced levels of HIF-1α after treatment with αMG

MCF-7 cell monolayers were exposed to 1% O₂ tension for 48 h and the ability of αMG to affect the intracellular levels of hypoxia-induced factor-1α (HIF-1α) was investigated. Under these conditions, it was possible to detect HIF-1α by western blot and to assess that 0.5 µg/ml and 1.0 µg/ml αMG encapsulated into NPs reduced its cellular concentrations in a dose-dependent manner (Figure 6), irrespective of the absence or presence of the thioaptamer. Similar effects were also obtained using free αMG at the concentrations of 0.5 µg/ml and 1.0 µg/ml (data not shown).

Discussion

In the present study, we evaluated the ability of NPs containing αMG to cause damage to MCF-7 cells grown in 3D as MCTSs. A robust derangement of spheroids was evidenced by the irregular shape of their edge accompanied by a parallel dissociation and decrease in density of the tumour bulk. The augmented distance between cells that was observed after treatment with αMG-loaded NPs did not occur only at the periphery of the MCTSs but also in the innermost layers, indicating that this effect of αMG was extended to the overall volume of the spheroid. The antitumor activity of αMG delivered by NPs was investigated using MCTSs growing in Matrigel or gelatine. Both gels are usually employed to simulate the ECM of MCTSs but with different analytical end-points for tumour aggressiveness investigations. Specifically, Matrigel provides a suitable support for tumour invasion assays, while gelatine is a better substrate to evaluate cell migration resembling cell dissemination from a solid microtumour or micrometastases [21].

In our recent study [3], we underlined that αMG was able to dissociate MCSTs generated by MDA-MB-231 breast cancer cells without increasing their tendency to migrate in a gelatine environment. The cells of these spheroids even reduced their motility when exposed to αMG, an antitumor effect that was also demonstrated in carcinomas deriving from other tissues [22, 23]. We also reported that MCF-7 cells of MCTSs could not migrate in gelatine-coated plates because of the low aggressiveness of this breast cancer cell line. In the present study, the spheroids cultured on Matrigel- or gelatine-coated plates significantly reduced their compactness after treatment with αMG-loaded NPs but did not show any sign of invasiveness. The lack of invadopodia conferring a characteristic “starburst” pattern to MCTSs was a further confirmation that the looser aggregation of cells was not paralleled by an increase in malignancy. Therefore, the detachment of cells from MCTSs caused by αMG-loaded NPs can be really considered a significant anticancer effect that is not accompanied by the risk of increased invasiveness or metastatic spreading.

The lowest concentration of αMG as a cargo of NPs that provoked a substantial reduction in spheroid...
density in the presence of gelatine was 0.1 µg/ml and the greatest disaggregation was obtained with 0.5 µg/ml αMG. It is worth noting that these doses of αMG delivered by NPs are about tenfold inferior to those provoking damages to MCF-7 MCTSs with free αMG. Moreover, αMG delivered by NPs dissociated MCTSs that were cultured in Matrigel or gelatine. Thus, these results suggest that: (i) NPs are suitable vehicles allowing αMG to damage tumours at very low concentrations; (ii) the αMG-dependent loss of contacts shown by MCF-7 MCTS cells involves the presence of a gel containing collagen fibres. The finding that αMG can inhibit the adhesion of MCF-7 cells to type I collagen [24] supports the hypothesis that some components of the ECM are involved in the dissociation of MCTSs induced by the xanthone. It has been also reported that an inadequate contact of collagen to integrins β represented an early step blunting the FAK/Akt/ERK cascade that, in turn, provoked the downregulation of the NF-kB-driven expression of MPP2 and MPP9 metalloproteinases [25]. This last effect is considered one important mechanism responsible for the reduced invasiveness of tumour cells that follows a treatment with αMG [25]. Moreover, Kopp et al. demonstrated that the capability of MCF-7 cells to form MCTSs was related to the activity of NF-kB [26]. Thus, we supposed that the anti-adhesive property shown by αMG-loaded NPs towards breast cancer MCTSs was the result of the weaker bond that MCF-7 cells established with collagen fibres inside the spheroid and the consequent lack of NF-kB activation. Similarly, Ivascu et al. showed that an anti-integrin β1 antibody was able to dissociate MDA-MB-231 spheroids exclusively when they were grown in the presence of 2.5% Matrigel [27]. It is presumably that also under our conditions both Matrigel and gelatine fibres contributed to the supporting network of the growing MCTSs and that αMG counteracted some cellular activity related to the presence of collagen. Moreover, Ivascu et al. demonstrated that the dissociation of MCTSs did not imply the loss of viability of tumour cells [27]. Therefore, the use of low doses of αMG-loaded NPs could be investigated as a non-toxic adjuvant drug for tumour chemotherapy. Indeed, the αMG-driven cell dissociation might facilitate both penetration and diffusion of conventional drugs into the tumour bulk.

When αMG was delivered by the NPs conjugated to the CD44 thioaptamer, a loss in cell contact of tumour spheroids did also occur, but MCTSs underwent a lower size in respect to those obtained with NPs not linked to the thioaptamer. This suggests that a number of cells died or slowed down their process of duplication. To explain this effect, it is important to underline that at least 60% of MCF-7 cells express CD44 [28] and that these cells, by sequestering via a very efficient receptor-mediated process the αMG-loaded NPs conjugated to the thioaptamer, should be subjected to a more severe antitumor activity.

Several molecular mechanisms underlying the αMG action have been described, including the inhibition of HIF-1α in pancreatic stellate cells [29]. In the present study, we also observed that αMG was able to downregulate the protein expression of HIF-1α in MCF-7 cell monolayers after hypoxia activation. Thus, as previously suggested [30], it is conceivable that αMG can reduce the number of cancer cells that have been adapted to hypoxia by blunting the expression of HIF-1α. It is interesting to note that breast epithelial CSCs, which are protected from drug toxicity in their hypoxic “niches”, largely express CD44 together with the epithelial-specific antigen (ESA), whilst they stain CD24+/low [31]. Consequently, NPs bearing αMG and targeting CD44+ cells could represent a drug able to target both CSCs and cells of the tumour bulk in breast carcinoma. In addition, the clinic-pathological and adverse prognostic implications of the expression of HIF-1α that have been demonstrated in different cancer types [32, 33] reinforce the expectation towards the antitumor efficacy of drugs targeting HIF-1α.

Another advantage of NPs conjugated to a thioaptamer, rather than a normal aptamer, should be their lower clearance due to the elevated resistance that the sulphur-adenine nucleotides offer against the nuclease attack [34]. Moreover, the PLGA core of the NPs used in the present study was covered by phospholipids and PEG that have been demonstrated to increase their in vivo lifespan [11].

In conclusion, these results underlines how great is the antitumor effect of αMG delivered by lipidic NPs, since very low concentrations of the xanthone as a cargo caused a strong disaggregation of MCF-7 MCTSs. When αMG was loaded in CD44 thioaptamer-tagged NPs, it was still responsible of spheroid dissociation, but the toxic effect was more relevant because accompanied by a reduction in MCTS size. Studies are in progress to ascertain whether the selectivity of αMG-loaded NPs linked to a CD44 thioaptamer may also represent a strategy to destroy CSCs in breast carcinomas.

**Abbreviations**

αMG: α-mangostin; b-FGF: basic fibroblast growth factor; DMEM: Dulbecco's modified Eagle medium; ECACC: The European Collection of Authenticated Cell Culture; EGF: epidermal growth factor; ESA: epithelial-specific antigen; MCTS:
multicellular tumour spheroid; ULA: ultra-low attachment.

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Authors’ contributions

F. Bonafe, C. Pazzini: concept/design/acquisition of data, data analysis/interpretation.
S. Marchionni: SEM analysis, critical revision of the manuscript.
C. Guarnieri: critical revision of the manuscript.
C. Muscari (principal investigator): concept/design/acquisition of data, data analysis/interpretation, drafting of the manuscript, critical revision of the manuscript.

Competing Interests

The authors have declared that no competing interest exists.

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