Silica nanoparticle surface chemistry: An important trait affecting cellular biocompatibility in two and three dimensional culture systems

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- PMSN resulted in disrupted morphology and very low viability in the 2D cell culture

Abstract

Great advantages bestowed by mesoporous silica nanoparticles (MSNs) including high surface area, tailorable pore diameter and surface chemistry, and large pore volume render them as efficient tools in biomedical applications. Herein, MSNs with different surface chemistries were synthesized and investigated in terms of biocompatibility and their impact on the morphology of bone marrow-derived mesenchymal stem cells both in 2D and 3D culture systems. Bare MSNs (BMSNs) were synthesized by template removing method using tetraethylorthosilicate (TEOS) as a precursor. The as-prepared BMSNs were then used to prepare amine-functionalized (AMSNs), carboxyl-functionalized (CMSNs) and polymeric amine-functionalized (PMSNs) samples, consecutively. These nanoparticles were characterized by scanning electron microscopy, zeta potential measurement, dynamic light scattering, BET (Brunauer, Emmett, Teller) analysis, and FTIR technique. In a 3D culture system, stem cells were encapsulated in alginate hydrogel in which MSNs of different functionalities were incorporated. The results showed good biocompatibility for both BMSNs and AMSNs in 2D and 3D culture systems. For these samples, the viability of about 80 % was acquired after 2 weeks of 3D culture. When compared to the control, CMSNs caused higher cell proliferation in the 2D culture; while they showed cytotoxic effects in the 3D culture system. Interestingly, polymeric amine-functionalized silica nanoparticles (PMSNs) resulted in disrupted morphology and very low viability in the 2D cell culture and even less viability in 3D environment in comparison to BMSNs and AMSNs. This significant decrease in cell viability was attributed to the striking uptake values of highly positively charged PMSNs by cells that were measured using inductively coupled plasma

optical emission spectroscopy instrument (ICP-OES). These results uncover different interactions between cell and nanoparticles with various surface chemistries. Building on these results, new windows are opened for employing biocompatible nanoparticles such as BMSNs and AMSNs, even at high concentrations, as potential cargos for carrying required growth and/or differentiation factors for tissue engineering applications.

Key words: Mesoporous silica nanoparticles, human mesenchymal stem cell, poly-L-arginine, alginate, cytotoxicity, tissue engineering

Highlights:

- Suitable biocompatibility of both BMSNs and AMSNs in 2D and 3D hMSCs culture systems
- CMSNs caused higher cell proliferation in the 2D culture system
- CMSNs showed cytotoxic effects in the 3D alginate-based culture system
- PMSNs resulted in disrupted morphology and very low viability in the 2D cell culture

1. Introduction

In the past few decades, nanomaterials have been cemented as promising tools in tissue engineering, drug delivery, and cancer therapy [1–4]. For example, nanomaterials such as graphene, carbon nanotubes, hydroxyapatite particles and nanodiamonds have shown great potential in bestowing superior electrical and mechanical properties to scaffolds required for engineering specific types of tissues [3,5–8]. Indeed, improved electrical conductivity, magnetic properties, and antimicrobial activity are some of the advantages of noble metallic and metal oxide nanostructures such as gold, silver, iron oxide, and titanium nanoparticles; which in turn makes them suitable for biomedical applications [9–11]. Since the discovery of bioactive inorganic materials in animal body, natural and synthetic inorganic nanoparticles have gained a great attention in tissue engineering. A wide range of these nanomaterials including

nanohydroxyapatite (nHA), silicate, bioactive glasses, and calcium phosphate nanoparticles, have been studied for musculoskeletal tissue engineering applications [12–15].

Among the many different types of inorganic nanomaterials, mesoporous silica nanoparticles (MSNs) hold the greatest promise in biomedical applications. Fig. 1 presents the number of articles published per year with a title containing “mesoporous silica nanoparticle” and the number of citing articles. The steady increase in these numbers indicates a growing interest to MSNs and their applications in different areas. This is certainly due to the attractive properties they present, such as high surface area and pore volume, which ensure sufficient loading and effective release of therapeutic agents [16–19]. Notably, their straightforward synthesis procedure, tunable pore sizes and facile surface functionalization as well as their great biocompatibility have made MSNs excellent candidates in many areas of biomedical applications [16,18,20]. As an example, various types of theranostic agents for cancer diagnosis and therapy were developed based on MSNs in recent years [21–24]. MSNs have also been employed to reinforce the mechanical properties of biomaterial scaffolds [25], and to provide biological cues required for directing stem cells toward desired lineages [26–28].

Fig. 1: Number of articles published with a title containing “mesoporous silica nanoparticle” shown in red bars. Number of publications citing the respected articles per year, shown in blue bars. The data were extracted from Web of Science, Clarivate

Analytics (July 2018).

One of the most important characteristics of biomaterials for tissue engineering is the way they interact with cultured cells. Among the many yet unrecognized interactions, the most basic one is the extent to which a nonnative matter can be applied without cytotoxic effect, namely the material biocompatibility [29]. These interactions between nanomaterials and cells can affect cells morphology, proliferation and fate. It has been shown that the different morphologies in the effect of different culture systems regulate hMSCs fate to either adipocyte or osteoblast cells [30]. On the other hand, the cells morphology can affect the nanoparticles uptake rate and consequently the degree of their cytotoxicity [31]. Farvadi et al. indicated that the cell uptake and cytotoxicity of gold nanorods with an average size of 33.5 ± 5.5 nm are strongly dependent on the morphology of the normal fibroblast and the cancerous epithelial (MCF-7) cells [31].

Our hypothesis is that decorating MSNs with different functional groups occur a great impact on biocompatibility of colloidal nanoparticles. In this research, we study and compare the biocompatibility of MSNs with different surface functionalities (amine, carboxyl and polyamine) in both 2D and 3D culture systems. Herein, mesoporous silica nanoparticles were synthesized by template removing method. Then, as illustrated in Fig. 1a, they were modified to affix different functional groups on their surface. The as-prepared MSNs were introduced to 2D and 3D cell culture systems in different concentrations and their effects on the cell morphology and viability in the culture systems were studied. A simple schematic of 2D and 3D cell cultures developed in this work are demonstrated in Fig. 1b.

Fig. 2: a) Schematic illustration of mesoporous silica nanoparticles, the precusures used for functionalization and their final functionalities, dicriminated in different colors. b) Schematic illustration of MSNs biocompatibility tests in 2D and 3D culture systems. In 2D culture system, hMSCs were cultured in media containing MSNs with different functionalities. To evaluate the the cytotoxicity effects in the 3D culture system, stem cells were encapsulated in an alginate hydrogel incorporated with MSNs of differnet functionalities.

2. Materials and methods

2.1. Materials

Alginate (Al, pharmaceutical grade, FMC Biopolymer, UK), tetraethylorthosilicate (TEOS, Aldrich-86578), hexadecyltrimethylammonium bromide (CTAB, Merck-814119, Germany), 3-aminopropyl triethoxysilane (APTES, Sigma-Aldrich-A3648), N,N-dimethylformamide (DMF,

Sigma-Aldrich-33120), poly(L-arginine hydrochloride) (MW = 19000, Alamanda polymers, USA), succinic anhydride (Sigma-Aldrich-239690), acetic acid glacial (Sigma-Aldrich-537020), N-Hydroxysuccinimide (NHS, Sigma-Aldrich-130672), N-(3-dimethylaminopropyl)-N-ethylcarbodiimide hydrochloride (EDC, Sigma-Aldrich-03450), ethanol absolute (VWR chemicals), hydrochloric acid (HCL, 37 %, Sigma-Aldrich-320331), nitric acid (Merck, Germany), sodium hydroxide (NaOH, Sigma-Aldrich-S0899), and Dulbecco's phosphate-buffered saline (DPBS, Thermo Fisher Scientific) were purchased from the above-noted companies and used as provided. Water was deionized using water purification system (Milli-Q Academic, Millipore, USA).

Human mesenchymal stem cells (hMSCs) were delivered by Lonza Inc. and preserved as freeze stock. Low glucose Dulbecco's modified Eagle's medium with GlutaMAX™ supplement (DMEM), fetal bovine serum (FBS), and penicillin/streptomycin were purchased from Thermo Fisher Scientific company. Abcam-ab112118 cell cytotoxicity kit were purchased from Abcam company. Nunc-MicroWell 96-well optical-bottom plates were bought from Thermo Fisher Scientific company. LIVE/DEAD® viability assay kit was purchased from Life Technologies and used according to the manufacturer protocol. Silicon isolators (JTR24R-1) were procured from Grace Bio-Labs.

2.2. MSNs synthesis

Template removing method was used to synthesize mesoporous silica nanoparticles [32]. Briefly, 100 mg of CTAB as liquid crystal templating agent was dissolved in 48.65 ml of deionized water. The pH was fine-tuned between 10.5 to 11.5 using sodium hydroxide solution (2 M). TEOS (1 ml) was added dropwise as the silica precursor while the solution was stirred (1500 rpm) at 80 °C. After 2 hours, the solution was centrifuged, and the MSNs were subsequently collected from

the sediment layer. In order to remove the remaining CTAB, the MSNs were washed two times for 8 hours in a 10 % (v/v) solution of hydrochloric acid in almost pure ethanol, followed by final washing with absolute ethanol and deionized water. They were then kept at 4 °C for further down-stream usages.

2.3. Surface functionalization of MSNs

3-(2-aminoethyl amino)propyltrimethoxysilane, succinic anhydride, and poly-L-arginine were used to functionalize MSNs surface with amine, carboxylic and polymeric amine groups, respectively. In order to prepare amine-functionalized MSNs, ethanol-washed MSNs (150 mg) were dispersed in 13.8 ml of absolute ethanol using sonication, then 0.6 ml of deionized water and 0.3 ml acetic acid was added. APTES (0.3 ml) was introduced to the solution at room temperature while being stirred at 1000 rpm. Once the reaction was completed, the as-prepared AMSNs were washed twice with ethanol and deionized water and stored in fridge for further use.

To prepare carboxyl-functionalized MSNs, succinic anhydride (200 mg) was added to a synthesis vessel containing DMF (20 ml) stirring at 1000 rpm under nitrogen purge. After 20 min, a suspension of well-dispersed AMSNs in DMF (20 ml, 0.5 % w/v) was added in a drop-wise manner during 10 min to the first solution while being stirred at 1000 rpm. After additional 20 minutes of nitrogen purging, the vessel was sealed with parafilm and left at room temperature for 24 hours. The CMSNs were washed 2 times with DMF, ethanol, and deionized water sequentially and stored in deionized water in fridge for further use.

Deionized water-washed CMSNs were used in order to prepare poly-L-arginine-coated MSNs (PMSNs). For that, CMSNs (50 mg) were dispersed in 10 ml of PBS solution (10 mM) containing 2 mg NHS and 10 mg EDC. Then, 1.5 ml aliquots of CMSNs suspension were shaken for 7 min. The nanoparticles were separated from each aliquot by centrifugation (3 min, 5000

rpm) and then, were dispersed in 1.5 ml of a previously prepared PBS solution (10 mM) containing poly-L-Arginine (0.2 % w/v) and were shaken for 1 hour. Finally, the prepared PMSNs were washed with PBS solution (10 mM) and deionized water 3 times (3 min, 5000 rpm) and stored in fridge for further use.

2.4. MSNs characterization

To determine the mean diameter of the prepared MSNs, samples were coated with gold (10 nm) and imaged using SEM (FEI-Quanta 200 ESEM FEG) operating at 10 kV. The presented mean diameter was estimated using Fiji, an image processing package for biological-image analysis, for more than 500 nanoparticles [33].

As CTAB is a biologically toxic material, it should be completely removed after the washing step explained in the previous section. In this regard, the samples were freeze-dried and CTAB removal was evaluated with Fourier transform infrared (ATR-FTIR) spectroscopy (PerkinElmer Spectrum 100 FTIR spectrometer, USA). The transmittance spectrum of the respective samples was recorded between 4000 to 600 cm^{-1} .

To validate the successful surface modification of MSNs, FTIR spectra of the prepared samples were recorded using Bruker tensor 27 FTIR spectrometer. To prepare the samples, all four types of MSNs were thoroughly dried on a hotplate at 120 °C for 2 h. One mg of each sample was used for preparing KBr pellets.

Zeta potential of MSNs functionalized with different chemical groups was measured at 25 °C using Malvern Zetasizer device (Nano Series ZS, Malvern Instruments, UK). The hydrodynamic size of MSNs was measured by dynamic light scattering (DLS) using Nano S device (Malvern, UK). To this end, the samples were dispersed in deionized water (0.1 mg/l) and sonicated for 15 min before running the test. BET (Brunauer, Emmett, Teller) analysis was employed using a surface

area analyzer (Micromeritics TriStar II 3020, USA) to determine the specific surface area of the synthesized MSNs. To confirm the mesoporous structure of MSNs, BJH (Barrett-Joyner-Halenda) method revealing MSNs' pore volume and size distribution was employed.

2.5. Cell culture

hMSCs were expanded in tissue culture flasks in DMEM supplemented with 10 % FBS and 1 % penicillin/streptomycin and were incubated in a humidified environment with 5 % CO₂ at 37 °C until further use. Cells were used at passage 3-5 for all cell toxicity and viability tests.

2.6. Cell encapsulation

The prepolymer solution containing MSNs of different concentrations was mixed with hMSCs to obtain the final concentration of 10⁶ cells/ml of hydrogel. Then, the prepolymer solution was deposited using an automated system (Eppendorf Xplorer, H31141F, Eppendorf, Germany) into silicon isolators (2 mm diameter × 1 mm depth) which previously sterilized and mounted on glass slides. To crosslink the polymeric chains of alginate with the final concentration of 1 % w/v and make the cell-laden hydrogels, each microwell was covered by 10-20 µL of a CaCl₂ solution (2 % w/v). After 5 minutes, to remove the excess salt, the as-prepared hydrogels were washed twice using DPBS and covered with enough complete medium. The samples were incubated in humidified environment with 5% CO₂ at 37 °C during which the medium was changed every 3-5 days.

2.7. Viability studies

To monitor the cytocompatibility of MSNs, the respected tests were performed in 2D and 3D cell culture systems. The cytotoxicity assays in 2D cell cultures were conducted using Abcam cytotoxicity kit as suggested in the manufacturer protocol. Briefly, cells were seeded at the concentration of 5 × 10⁴ cells/ml in 96-well plates and were incubated for one day. Then, the

media were replaced with 100 μ l of fresh media containing the respected amount of MSNs. After 2 days of incubation, 20 μ l of the assay solution was added to each well and the samples were incubated for one more day. The media (100 μ l) were transferred to new well plates and the absorbance was recorded at 570 nm and 605 nm. The cell viability was then calculated according to the following equation:

$$\text{Cell viability (\%)} = (R_{\text{Sample}} - R_o) / (R_{\text{Control}} - R_o) \times 100 \quad (1)$$

here, R_{Sample} is the absorbance ratio (OD_{570}/OD_{605}) of samples cultured with MSNs. R_{Control} is the absorbance ratio (OD_{570}/OD_{605}) for the samples cultured in normal media. R_o is the average of the absorbance ratio of the control samples containing no cells.

Fluorescent staining and imaging were applied to determine how MSNs concentration and surface chemistry affect hMSCs morphology and viability. hMSCs were seeded in a 48-well plate (2×10^4 per well) by dispensing 250 μ l media and incubated for one day. The old media were changed with the media containing the respected nanoparticles. After 48 h of incubation, media were removed and 70 μ l of previously prepared LIVE/DEAD® viability assay solution was added to each well. Using this assay, live and dead cells are stained with calcein-AM (green) and ethidium homodimer-1 (red), respectively. Then, a confocal microscope (CLSM, Zeiss LSM 700, Germany) was employed to image the stained cells.

To investigate the biocompatibility of the hydrogel nanocomposite in 3D culture system, the encapsulated cells were stained after being cultured for 1, 3, 7, and 14 days using LIVE/DEAD® viability assay kit and the suggested protocol by the manufacturer. The live and dead cells were imaged using a confocal laser-scanning microscope and the images were analyzed by CellProfiler software. To this end, the red and green channels were first separated using the built-in modules in the software, and the cells in each color channel (green for live cells and red for dead cells)

were identified and counted. Finally, the following equation was used to evaluate the cell viability:

$$\text{Viability (\%)} = \frac{\text{Number of live cells}}{\text{Total cell number}} \times 100 \quad (5)$$

2.8. Cellular uptake of MSNs

To measure cellular uptake of MSNs, hMSCs were seeded at 2×10^5 cells/flask in T25 flasks. After one day, the cells were treated with MSNs of different surface chemistries (100 $\mu\text{g/ml}$) for two days. As the control sample, the same number of cells were cultured in a flask with no treatment. To remove any free MSNs, media was removed and the cells were washed with PBS twice; after which the cells were harvested by trypsinization and centrifugation. The precipitated cells were resuspended in PBS for cell counting. Then, the pellet was dissolved in 500 μl of concentrated nitric acid (100 % v/v). Afterward, the final volume of each sample reached to 4 ml using DI water and was incubated at 50 $^{\circ}\text{C}$ for 30 min. The samples were stored in the fridge until their introduction to inductively coupled plasma optical emission spectroscopy instrument (ICP-OES, Arcos, Spectro, Germany) for detection of Si internalization by cells.

2.9. Statistical analysis

To perform statistical analysis of the acquired data GraphPad Prism 7.01 software was employed (San Diego, USA). All numerical results were reported as mean $\pm$ standard deviation and Gaussian distribution was evaluated using Shapiro–Wilk normality test. Using Brown–Forsythe and Bartlett’s tests, homogeneity of variances was evaluated, and for data groups in compliance with the prerequisites of one-way ANOVA, the significance of difference was examined through one-way ANOVA and Tukey’s post hoc test. Otherwise, the Kruskal–Wallis nonparametric test followed by Dunn’s multiple comparison test was performed. The statistical

significance levels were presented as $^*(p<0.05)$, $^{**}(p<0.01)$, $^{***}(p<0.001)$, and $^{****}(p<0.0001)$.

3. Results and discussion

3.1. MSNs characterization

Images taken by scanning electron microscopy (SEM) indicated that the synthesized MSNs featured spherical shape with uniform diameter (Fig. 3a). To measure the mean diameter of MSNs and investigate their monodispersity, Fiji analysis program was employed (Fig. 3b). The acquired results showed a mean diameter of 88 ± 13 nm with a PDI value of 0.15 implying that the synthesized MSNs are monodisperse and have a size range that enables a high cellular uptake [34]. From the FTIR results (Fig. 3c), we noticed that the two intense peaks at 2854 cm^{-1} and 2924 cm^{-1} which represent the symmetric and asymmetric stretching vibrations of the methylene chains of CTAB vanish completely through ethanol-HCl treatment [35]. This implies that the synthesized MSNs are free of cytotoxic precursors.

An important characteristic of nanoparticles is their surface charge; which can significantly influence their interactions and uptake by cells as well as their cytocompatibility [36–38]. Because of the innate negative charge of the cells membrane, positively charged nanoparticles result in higher cellular uptake making them more efficient carriers for delivery of biomolecules [38]. Nevertheless, when compared to their low charged counterparts, the higher cytotoxicity of highly-charged polycationic constructs has previously been reported [37]. Fig. 4a shows the zeta (ζ) potential values of the MSNs with different surface chemistries. The ζ value of -18 mV for BMSNs is attributed to presence of hydroxyl groups on their surface. Suggesting successful surface functionalization of BMSNs, this value was changed to about +26, -20, +43 mV once modified with amine, carboxyl, and polyamine groups, respectively.

Fig. 3: a) SEM micrograph showing mesoporous silica nanoparticles. b) Size distribution of synthesized silica nanoparticles. c) FT-IR spectra showing the elimination of methyle, methylen, and amine groups related to CTAB removal from the as-prepared MSNs by treating with ethanol-HCl solution.

Furthermore, DLS analysis was employed to explore the effect of the surface chemistry on the dispersion/agglomeration of MSNs. Fig. 4b shows the MSNs diameter distribution on which regression of gaussian equation was applied. DLS measures the hydrodynamic diameter of MSNs, which is strongly dependent on the solvent and its layers associated with MSNs while in microscopy techniques, the actual size of MSNs is measured in its dry state [39]. The hydrodynamic diameters of BMSNs, AMSNs, and CMSNs were obtained 240 ± 27 , 276 ± 31 , and 199 ± 22 nm, respectively (Fig. 4b). The significant difference in the hydrodynamic size of PMSNs' diameter (103 ± 10 nm) in comparison to other MSNs might be attributed to its high zeta potential value (+43 mV) which supports their good dispersion and improves the PMSNs colloidal stability.

Fig. 4c depicts FTIR spectra for the as-prepared MSNs. Siloxanes as the main groups in silica nanoparticles were characterized by stretching frequencies of Si-O-Si linkage in the range of $1100\text{--}1000\text{ cm}^{-1}$ [40]. Silanol (Si-OH) groups existing on BMSNs surface involve the hydroxyl (OH) groups stretch at $3700\text{--}3200\text{ cm}^{-1}$ [40–42]. The weak band at 1410 cm^{-1} confirmed the Si-C bond in all the surface modified MSNs, as described previously [40], which is a result of its functionalization with APTES. As OH moieties on silica surface were decorated by amine groups,

the width of its respected broad band ($3700\text{--}3200\text{ cm}^{-1}$) was reduced and replaced with amine (NH_2) group stretching starting from 3300 cm^{-1} [40]. The observed weak signal at 2940 cm^{-1} corresponding to stretching vibration modes of aliphatic C–H bonds was emerged after amine-functionalization of BMSNs and intensified after consequential functionalization [42]. In response to MSNs modification with carboxylic acids, a very broad band related to OH groups starting from 3200 cm^{-1} was shown up. The presence of carboxylic groups was also confirmed by a sharp band at 1720 cm^{-1} corresponding to the antisymmetric stretch of carbonyl groups in CMSNs [40]. Furthermore, the peak at 2979 cm^{-1} was related to OH groups in carboxylic acid dimers [40]. Ultimately, the successful decoration of MSNs by poly-L-arginine was validated by peaks at 1642 cm^{-1} and 1555 cm^{-1} corresponding to amide I and amide II, respectively [40]. The weak stretching band of NH also was appeared in the region of $3325\text{--}3500\text{ cm}^{-1}$ [40].

Fig. 4: a) Zeta potential of silica nanoparticles in aqueous solution confirming the favored functionalization. b) Hydrodynamic diameter of different MSNs acquired from dynamic light scattering. c) FT-IR spectra of MSNs of different surface chemistries (BMSNs, AMSNs, CMSNs and PMSNs).

In BET analysis, nitrogen isotherms of MSNs showed a great difference following various surface modifications. As results presented in table 1 show, the BET surface area and the total pore volume are dramatically decreased after consecutive modifications. For example, the BET

surface area decreased from 734 m²/g for BMSNs to 134 m²/g for PMSNs which confirms the successful binding of polymeric groups to both MSNs' internal and external surfaces.

Table 1. Structural properties of the synthesized MSNs

Nanoparticle type	$S_{\text{BET}}^{\text{a}}$ (m ² /g)	V_{t}^{b} (cm ³ /g)	$d_{\text{BJH}}^{\text{c}}$ (nm)	ζ^{d} (mV)	$d_{\text{DLS}}^{\text{e}}$ (nm)
BMSNs	734	0.713	2.5	-18	240 ± 27
AMSNs	579	0.585	2.3	+26	276 ± 31
CMSNs	486	0.461	2.1	-20	199 ± 22 nm
PMSNs	134	0.179	0.9	+43	103 ± 10 nm

a Total surface area from BET analysis
b Total pore volume calculated from nitrogen isotherm at $P/P_0 = 0.98$
c BJH adsorption average pore width
d Zeta potential
e Hydrodynamic diameter

3.2. Cytotoxicity and cell morphology

In order to determine the nanoparticle toxicity on 2D cultured hMSCs, a calorimetric kit was used to assay the viability by measuring the absorption ratio – a number that is directly proportional to the number of living cells. The presented results in Fig. 5a indicated that there are no cytotoxic effects observed in the MSNs concentration of 10 µg/ml. As the concentration increases to 50 µg/ml, the MSNs with the highest positive charge (PMSNs) showed significant cytotoxicity effects. In contrast, there was no cytotoxicity effect observed for BMSNs and CMSNs, even at the highest employed concentrations (1000 µg/ml). Meanwhile, no cytotoxicity occurred when applying AMSNs in concentrations up to 500 µg/ml. Therefore, AMSNs are suitable nano-carriers considering their non-cytotoxic property at relatively high concentrations and also a positive surface charge which result in higher cellular uptake and more efficient delivery of biomolecules [37].

To investigate in more detail what impact MSNs with different surface chemistries have

on cell toxicity, hMSCs treated with different concentrations of MSNs were stained using calcein-AM and ethidium homodimer-1 (Fig. 5b-f). Calcein-AM unravels the morphology of the living cells; while, ethidium homodimer-1 stains the nuclei of the dead cells. The fluorescent images showed almost no changes on the morphology of hMSCs treated even with the highest concentrations of BMSNs, AMSNs, and CMSNs (Fig. 5c-e) in comparison to control sample (Fig. 5b). However, it was observed that PMSNs disrupt the plasma-membrane integrity and spindle-shape morphology of the hMSCs (Fig. 5f). It is reported that surpassing a threshold of surface charge density may cause acute disruption of the membrane and therefore an intense cytotoxicity [43]. Poly-L-arginine may also, similar to polyethylene imine, cause enhanced cell uptake and subsequently destabilization of lysosomes within cells resulting in their content discharge in cytosol and ultimately cell apoptosis [44–47]. To further delve on these hypotheses, the effect of the different surface chemistries on MSNs uptake by hMSCs was assessed using ICP-OES technique. The obtained results revealed that hMSCs uptake PMSNs significantly (29.6 pg/cell) more than that of the BMSNs (1 pg/cell), AMSNs (2.3 pg/cell) and CMSNs (0.9 pg/cell); that most likely caused a 50% reduction in the cell viability by PMSNs at 100 $\mu\text{g/ml}$. Although PMSNs were observed to be highly cytotoxic at high concentrations, they showed great biocompatibility at 10 $\mu\text{g/ml}$. Therefore, this concentration might be applicable in controlled release delivery systems considering that the polymeric chains on the PMSNs surface may provide higher loading capacity than their counterparts. Moreover, L-arginine groups were reported to bestow many therapeutic benefits ranging from cancer immunotherapy [48] and anti-inflammation [49] to homeostasis [50,51].

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3.3. Cell viability in 3D environment

As the native cells in the body live in a 3D microenvironment and their phenotype and expression of transcription factors are completely different from that in 2D culture conditions [52,53], the viability of hMSCs encapsulated in alginate nanocomposite hydrogels containing different percentage of MSNs was studied. Fig. 6a-b depicts the live and dead cells after 2 weeks of culture in control (alginate) and nanocomposite hydrogels. Fig. 6c presents the results obtained on the hMSCs viability after 1, 3, 7, and 14 days cultures in different nanocomposite hydrogels using analysis of fluorescent images. These results indicate that the cell viability was almost over 70% for all the combinations after 7 days of culture. This high viability of hMSCs was maintained after 2 weeks for all the combinations except the hydrogels incorporated with more than 5% of CMSNs and PMSNs. This shows higher cytotoxic effect of CMSNs in 3D microenvironment when compared to 2D condition, which might be attributed to the different morphology of the cells in different culture systems, imposed by microenvironmental differences cells would sense in these two systems [31]. The spherical morphology of the hMSCs in 3D culture system (Fig. 6a-b) may imply unsuitable conditions in the microenvironments of the cultured cells in hydrogels incorporated with 5% of CMSNs and PMSNs, as described in the literature [54]. The more extensive impact of high concentrations of MSNs, in particular for CMSNs in 3D culture in comparison to that in 2D culture, can be due to this fact that the cells surface is exposed more to nanoparticles in 3D culture than in 2D culture. This can provide a higher chance for the interaction between cells and nanoparticles and consequently result in stronger impact on the cells in the 3D culture system. In this regard, future studies investigating cellular uptake for MSNs with different surface chemistries in 3D cultures and exact mechanisms of different cytotoxicity responses in different culture systems are of high importance. Moreover, there are some studies

reporting the differentiation-directing potential of small functional groups existing in the cell microenvironment [55–57]. In a study by Benoit et al. [57], for instance, a set of hydrophilic and hydrophobic functional groups inserted in a PEG-based matrix was used to induce differentiation of hMSCs toward osteogenesis, chondrogenesis, and adipogenesis. It has been shown that carboxylic groups mimicking the functional groups of glycosaminoglycans, the main part of native cartilage, upregulated expression of collagen II; whereas the hydrophobic moieties improved adipogenesis and amine groups showed no considerable changes [57]. Therefore, future studies assessing this potential of MSNs are highly suggested in order to further highlight the prospective application of the functionalized MSNs towards directing the stem cells differentiation.

Fig. 6: Live (green) and dead (red) cells visualized by confocal fluorescent microscopy after 2 weeks of culture in (a) nanocomposite hydrogels comprising different percentage of MSNs of various surface chemistries and (b) pure alginate hydrogel. c) The results of viability assay for hMSCs encapsulated in the alginate-based nanocomposite hydrogels after 1, 3, 7, and 14 days culture.

4. Conflicts of interest

There are no conflicts of interest to declare.

5. Conclusion

Herein, we have demonstrated the viability of hMSCs cultured in 2D and 3D systems in

the presence of MSNs with various surface chemistries. BMSNs and AMSNs showed no significant toxicity in the different concentrations tested in this study on hMSCs encapsulated in alginate hydrogels. In contrary, CMSNs and PMSNs showed significant cytotoxic effects at concentrations higher than 5% in such 3D cultures. It was demonstrated that the cellular uptake of PMSNs was greatly higher in comparison to those of other MSNs as a result of considerably high positive charge of PMSNs. That caused a significant decrease in the viability of cells incubated with even low concentrations of PMSNs.

Further studies are needed to investigate in detail the mechanisms governing the effect of nanoparticles on living cells, in particular in 3D cell culture systems. The high viability of the hMSCs cultured along with bare and amine-modified MSNs (BMSNs and AMSNS, respectively) in both 2D and 3D systems presents the potential of such nanoparticles as promising carriers of different types of biomolecules, such as growth and differentiation factors for a wide range of biomedical applications.

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