



Recent advances in stimuli-responsive drug release and targeting concepts using mesoporous silica nanoparticles

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Abstract

Being a developed and promising approach, nanotechnology has attracted a lot of attention in biomedical and pharmaceutical therapy applications. Among nanostructured materials, mesoporous silica nanoparticles (MSNs) are effectively used as nanocarriers for drug delivery systems. MSNs can be tailored-designed by different synthetic techniques. Their morphological characteristics dictate the type of application of such materials. Recently, polymer-based materials have been employed to functionalize the MSNs surface. These modified nanocarriers are loaded with the drug and can unload their “cargo” upon exposure to either endogenous or exogenous types of stimuli. In this study, different targeting concepts, including passive, active, vascular, nuclear, and multistage targeting, are discussed.

Keywords Mesoporous silica nanoparticles · Drug delivery · Endogenous and exogenous stimuli · Anticancer drug delivery · Targeting

1 Introduction

Drug delivery relies on the presence of a “neutral” carrier that allows for a “load” drug to reach the desired target and release the cargo at the site. This approach enables remarkably reduced side-effects from the drug effect at “off-target” regions as well as protecting the drug from degradation or solubilizing prematurely. Well-designed and functionalized nanoparticles have proven to be excellent vehicles to carry this task [1]. Another critical factor in this process is the ability of the drug to be released upon exerting a stimulus upon the carrier nanoparticles to unload its cargo at the desired site. In this respect, stimuli-responsive polymers were introduced to carry on this task [2]. Important characteristics have to be realized in these polymeric systems such as receiving, transmitting the stimulus, and responding conveniently and efficiently. Two types of stimuli triggers are identified: those are endogenous and exogenous types. Endogenous biological triggers include pH,

redox, glucose, and enzymatic systems. Endogenous type triggers anticipate the differences between the infected tissue and the microscopic environment surrounding this tissue in comparison to normal tissue physiology. Exogenous or external stimuli are, for instance, thermal, magnetic, and ultrasound effects. The latter have higher clinical applications because of the possible control over the carrier accumulation within certain positions around the targeted tissues and the precision in timing of the release of the load.

In the following sections, we will illustrate the synthetic strategies of mesoporous silica nanoparticles (MSNs) for different applications in drug delivery systems. The various techniques for applying exogenous and endogenous stimuli over MSNs for drug delivery will also be reviewed.

2 Synthesis of mesoporous silica materials

Mesoporous materials are defined by the international union of pure and applied chemistry (IUPAC) as those with pore size ranging between 2 and 50 nm arranged in an ordered pattern [3]. The first introduced mesoporous silica materials to the market were credited to those synthesized by Mobil Research and Development Corporation [4]. Those were designated commercial names as Mobil Crystalline Materials of Mobil Composition of Matter with an acronym of MCMs.

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The most popular ones are MCM-41 and MCM-48 and were introduced as catalysts. Extensive research was carried on these materials with efforts directed towards fine-tuning the structure of the resulting materials in terms of pore size. For example, MCM-48 nanoparticles (90 nm) demonstrated high permeability of encapsulated drug (resveratrol) compared to 150 and 300 nm ones [5]. On the other hand, the loading capacity of glucagon-like peptide-1 in dendritic silica nanoparticles (DSNPs) (pore size of 10 nm) was much higher than MCM-41 with small pores [6]. The employment of surfactants with designated chain lengths and structures are the key players in controlling mesoporous characteristics. For instance, MCM-41 has been widely studied as a potential candidate for drug delivery applications and was synthesized using cationic surfactants as intercalation templates [4, 7]. Moreover, cationic surfactant (CTAB) was used as a structure-directing agent for designing a biphasic system composed of MSNs and calcium phosphate for bone-specific drug delivery (doxycycline hydrochloride) [8]. The researchers in Mobil used alkyl trimethyl ammonium halide-based surfactants as templates and tetra-ethoxy silicate (TEOS)-sodium silicate-silica-Ludox mixture as silicate source. Besides, MCM-41 had a hexagonal structure; MCM-48 had a cubic structure while MCM-50 had a lamella structure. Several synthesis conditions affected the properties of the resulting mesoporous materials. These include the concentration of precursors, the type of solvent/co-solvent used, the temperature and pH of the synthesis solution, the calcination temperature, the chain length of the surfactant used, electrolytes/solubilizing agents, and co-surfactants incorporation to the synthesis solution [7, 9–17].

Another category of mesoporous silica materials widely used in biomedical applications was synthesized by a group of scientists at the University of California, Santa Barbara, designated as SBA [18]. Relatively larger pores characterize SBA-based silica mesoporous materials compared to those of MCM- materials. The synthesis of SBA-materials relied on the use of non-ionic tri-block copolymers such as alkyl poly (ethylene oxide) and oligomer surfactants. The resulting mesoporous materials are SBA-11 with cubic structure, SBA-12 with a 3D hexagonal shape, SBA-15 with a hexagonal shape, and SBA-16 with cubic caged-structure. The designated form depends on the symmetry of the mesoporous structure and the polymers used [18].

Several mechanisms have been proposed for the preparation of these materials. Two mechanisms were initially suggested. Firstly, the silica framework is formed around pre-existing liquid crystalline phases. Secondly, the addition of inorganic species affected the surfactant assembly and resulted in ordered arrays [19]. However, the first mechanism was not well-established because of the surfactant amount used in the synthesis solution is short from realizing the formation of liquid crystalline phase in aqueous media. Several other

mechanisms were also cited, for instance, a mechanism described as “swelling-shrinking mechanism” was proposed and confirmed by time-resolved synchrotron small-angle X-ray scattering (SAXS) technique [20]. This mechanism is only valid when TEOS is the only precursor used and in the absence of any other ethanol-like solvent. Figure 1 showed the mechanism of formation of MCM-41 [22]. Figure 2 illustrated the mechanism of mesoporous silica nanoparticles (MSNs) according to the swelling-shrinking mechanism [22]. Comprehensive information can be found in the recently published review on the diverse routes of MSNs synthesis [23, 24]. Recently, novel virus-like MSNs consisted of nanospheres and peripheral perpendicular nanotubes were fabricated via a single micelle templating epitaxial growth strategy. These virus-like MSNs demonstrated superior cellular uptake property, good internalization ability, and prolonged blood circulation ($t_{1/2} = 2.16$ h) [25].

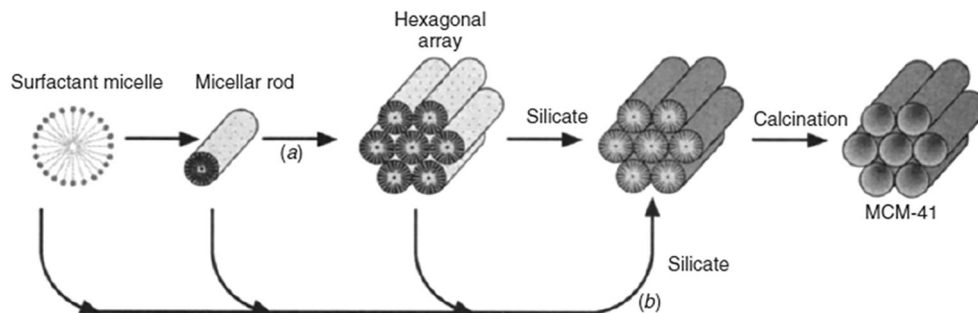
The more likely properties of MSNs that tailor the biomedical performance in terms of drug or gene loading and their release are particle size, pore size, and volume. For instance, relatively smaller particle sizes were obtained using some reagents, including alcohols, amines, inorganic bases, and inorganic salts [26–28]. Those agents affect the kinetics of hydrolysis and condensation reactions of the silica precursor. In 2014, a new generation of floating tablet-based MSNs was synthesized to enhance the delivery pattern of both hydrophobic (curcumin) and hydrophilic (captopril) drugs. The resultant tablets showed improved solubility for curcumin and prolonged release for captopril [29].

The pore size of the particles varied according to the chain length of the surfactants used. Thus, the applied surfactants with longer chain length resulted in the formation of larger pores [30]. Pore volumes, on the other hand, depended on the concentration of precursors and surfactants used. For example, the concentration of TEOS influenced the mesostructure ordering of the prepared MSNs [31]. Larger contents of TEOS resulted in more disordering in the mesostructure of particles, and insufficient contents hindered the mesoporous formation. The influence of the surfactant amount was discussed using cetyl ammonium bromide (CTAB) [21]. When CTAB was used with low concentrations, no micelles are formed, and no templates are available for the particle formation. Whereas, relatively high concentrations of CTAB resulted in disordered morphologies. Therefore, “moderate” concentrations of the surfactants are recommended for obtaining the desired mesoporous structure.

3 Stimulus-responsive gatekeeping of functionalized MSNs

Recently, designing a smart nanoparticle drug delivery system that responds to specific stimuli has gained a lot of attention

Fig. 1 Mechanism of formation of mesostructured surfactant-templated inorganic materials for the synthesis of MCM-alike materials: (a) Liquid-crystal phase initiation; (b) Silicate anion initiation. (reprinted with permission from Elder [21])



due to its tremendous promises in diagnostic as well as therapeutic application [32]. Two strategies are adopted in the synthesis of stimuli-responsive MSNs. In one approach, endogenous stimuli which are explicitly used for diseased tissue and proved to improve the drug action selectivity. It needs the addition of a suitable material to MSNs that respond to particular endogenous stimuli. The MSNs spontaneously enable the tumor-therapeutic selectivity and efficiency with bioresponsive therapeutic efficacy [33]. In the other strategy, it depends mainly on applying external physical stimuli to the desired tissue after administration of drug-loaded MSNs. Similarly, these applied exogenous stimuli lead to alteration of the MSNs structure and release of drug to the desired tissue [34]. Thus, researchers’ effort is aimed towards applying a combination of several stimuli-responsive strategies to enhance targeting efficiency and avoid the premature drug release [34, 35].

3.1 Exogenous stimuli-responsive drug delivery

Exogenous-type stimuli are applicable by using an external factor such as temperature, electromagnetic and magnetic fields, and ultra-sound. Endogenous or exogenous stimuli are aimed to trigger the release of a given drug desirably at targeted locations inside the body. This approach is advantageous in targeting the specific delivery of a particular drug to

its target site and limiting its side effects. The schematic diagram of Fig. 3 summarizes various exogenous and endogenous stimuli for drug delivery [36].

3.1.1 Thermal-triggered drug delivery/release

The strategy of this approach depends on changing the surrounding temperature around the particles loaded with the drug that resulted in triggering the pores around the surface of particles. Thermo-responsive drug delivery systems were widely implemented for anti-tumor treatment regimens [37]. This approach relies on coating or incorporating the MSNs with a thermo-responsive agent Fig. 4.

For example, MSNs were coated with organic/inorganic thermosensitive nanomaterials using reversible addition fragmentation transfer (RAFT) polymerization and click reaction [38]. Figure 5 demonstrates the coating of MSNs with thermo-responsive poly(N-isopropyl acrylamide) (PNIPAM). In this mechanism, the drug is released when subjected to heating stimulus. Similar approaches could be applied to combine two exogenous stimuli, such as magnetic field and thermal change. A recent study by Rinaldi et al. suggested the use of super-magnetic iron oxide nanoparticles that release heat as a response to an alternating magnetic field. Consequently, molecular loading was released via Diels-Alder bond scission [39]. The nanocarriers were coated with poly (ethylene

Fig. 2 Schematic of the structural changes of CTAB micelles before and after TEOS addition. (1) Before the addition of TEOS; (2) After the addition of TEOS; (3) The silica monomers are progressively released into the aqueous solution; (4) Silica monomers gradually release from the core of micelles and form a silica shell around micelles; (5) Silica forms a strong framework, and particles and mesopores are formed. (Reprinted with permission from reference 20)

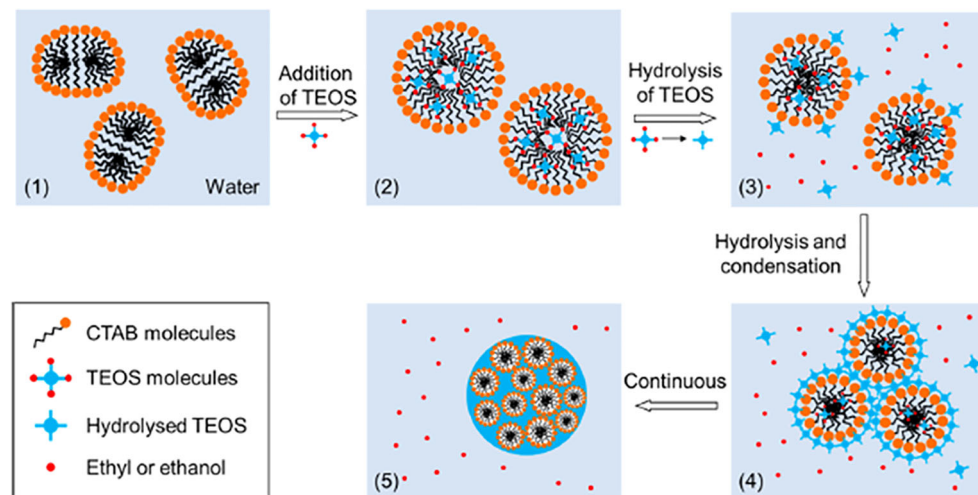
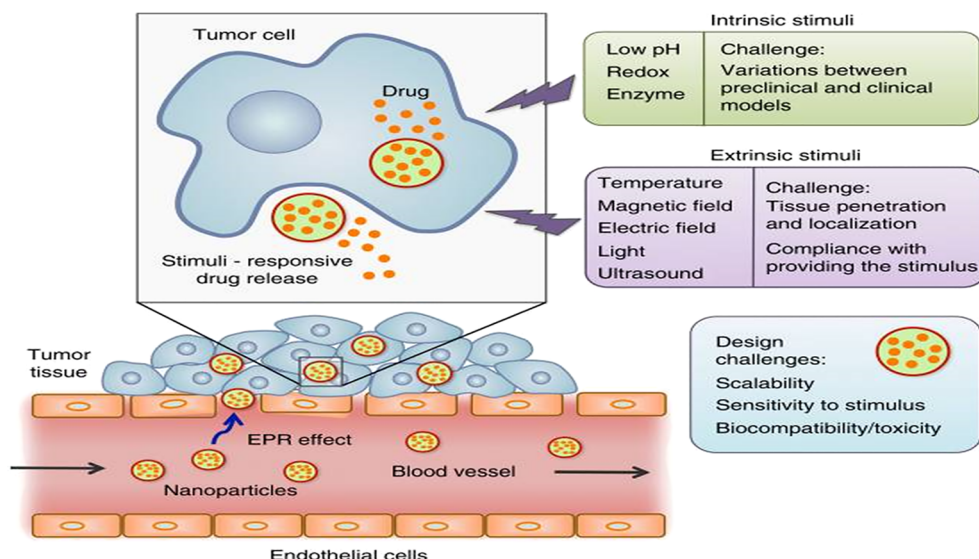


Fig. 3 Schematic diagram summarizing the various exogenous and endogenous stimuli for drug delivery (reprinted with permission from 36)



glycol)/poly (lactic acid) block copolymers. In their study, the authors reported that the release could be triggered multiple times using a magnetic field gradient to control the targeted regions for directive drug release spatially.

3.1.2 Magnetic-triggered drug release

Drug delivery based on magnetic trigger relies on releasing the magnetic and drug nanoparticles in response to the applied magnetic field. This performance ensures the directed delivery of magnetic nanoparticles to the targeted area in addition to controlling the drug release. Drug release in the response of the magnetic field relies on the ability of magnetic nanoparticles to produce thermal energy that induces morphological changes [40]. It was assumed that the magnetic core behaves like a hot spot that releases the drug without the need to elevate the global temperature [41] Fig. 6.

Iron oxide nanomaterials such as superparamagnetic nanoparticles were used extensively, replacing the MRI contrast agents [42]. Iron oxide nanoparticles have been linked to several ligands such as antibodies [43, 44], vitamins [45, 46], and polysaccharides [47]. In addition, a cyclic nine-amino acids peptide (Cys-Gly-Asn-Lys-Arg-Thr-Arg-Gly-Cys) (LyP-1) represented an excellent candidate for breast cancer, hepatocarcinoma, and atherosclerotic plaque therapies [46, 48]. The targeting of pancreatic cancer cells in vitro for diagnostic purposes was enhanced by using superparamagnetic mesoporous nanospheres anchored with lyp-1 ligand [49]. Doxorubicin has been loaded on graphene quantum dots (GQDs)-capped magnetic MSNs. These materials exhibited a synergistic effect against breast cancer 4T1 cells compared to chemo-hyperthermia and chemo-photothermal therapies. Under a magnetic field, the magnetic MSN/GQDs displayed heat liberation to the hyperthermia temperature [50]. Moreover, magnetic mesoporous silica microspheres with core-shell structure

Fig. 4 Schematic illustration of drug release from responsive MSNs triggered by temperature as an external stimulus

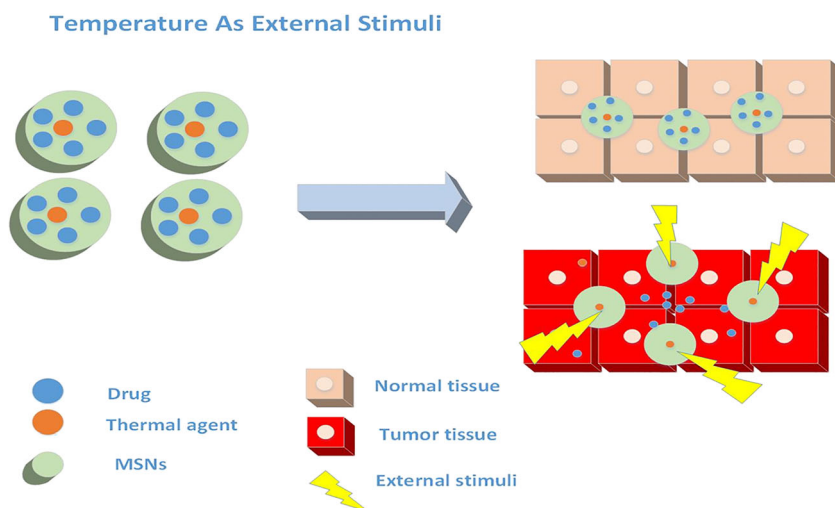
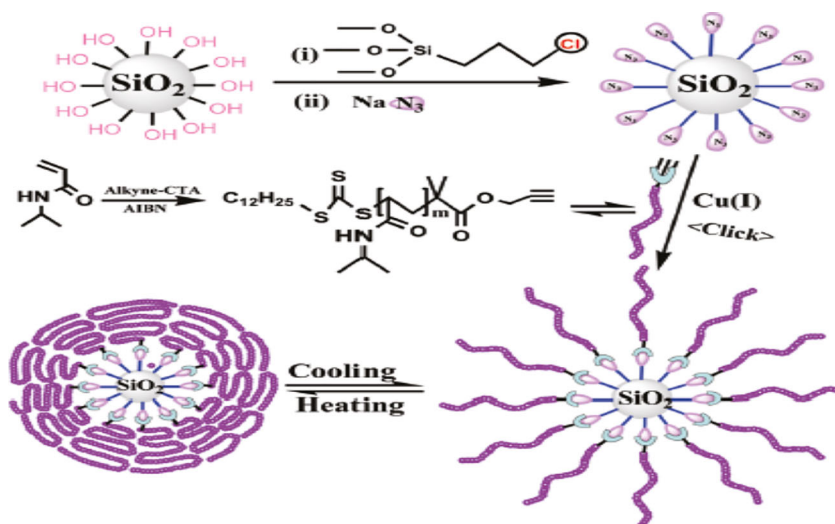


Fig. 5 Schematic diagram. Schematic illustrating the synthesis of MSNs coated with thermo-responsive PNIPAM brushes using RAFT polymerization and click chemistry (reprinted with permission from 38)



and large pores can be applied in bio-catalysis, bio-separation, and adsorption [51]. Surprisingly, Janßen et al. demonstrated that magnetic MSNs carriers did not accumulate on the magnetizable implant and area in orthopedics [52]. But instead, they were uptake and trapped in the lung, liver, and spleen of the mice with no significant pathomorphological alterations.

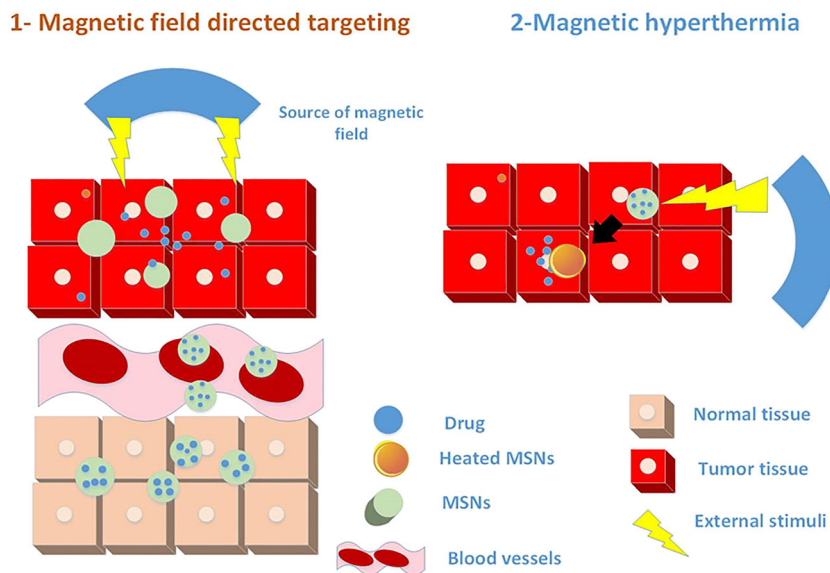
3.1.3 Light-triggered drug release

Activation delivery by light is a promising strategy due to its noninvasiveness property and remote control. It does not depend on changes in the chemical environment. The mechanism relies on conformational changes of the nanocarriers in response to light. Many light-triggered nanocarrier designs depend on releasing the drug from its cargo upon excitation by visible light or ultraviolet light. These nanocarriers proved

to be suitable only for vitro studies owing to its fast reduction in density [53]. Previous reports demonstrated that near-infrared gives maximum penetration, minimum absorbance, and more safety for tissues and organs than ultraviolet-visible light [54, 55]. Yang et al. fabricated smart and novel light stimulus MSN nanorods (MSNRs) doped with photosensitizer chlorin e6 (Ce6) for anticancer applications [56]. Doxorubicin release was achieved by coating MSNRs using bovine serum albumin through singlet oxygen-sensitive bis (alkylthio) alkene Bata linker (CMSNRs) and finally modified with polyethylene glycol. This carrier was activated via 660 nm with a low power density as low as 5 mW cm⁻² through SO-mediated cleavage of the Bata linker. In addition, they showed marked synergistic therapeutic effects in both in vivo and in vitro experiments.

Photodynamic therapy is an important therapeutic modality with high selectivity to the tumor and low invasiveness. A

Fig. 6 Schematic illustration of the mechanisms of drug release from MSNs triggered by a magnetic field and magnetic hyperthermia



light source activates the tumor localized photosensitizer, and this property results in producing highly toxic ROS such as singlet oxygen that has proven to suppress the tumor with high efficiency and specificity with little side effects [57]. In 2015, Liu et al. synthesized double mesoporous silica-shelled UCNPs that can deliver both photosensitizer and bio-reductive prodrug tirapazamine [58]. Upon excitation by laser at 980 nm, a large amount of ROS was produced that eradicate the tumor and show the feasibility of NIR-triggered deep PDT by MSN-based therapeutic. Multifunctional nanoplatform-based MSN-coated gold nanorods were fabricated for PDT/PTT synergistic therapy by Liu et al. [59]. MSN-based nanosensors were fabricated by incorporating a mesoporous silica shell with UCNP and oxygen indicator tris(4,7-diphenyl-1,10-phenanthroline) ruthenium (II) dichloride [60]. The mesoporous silica particles provide a large surface area that guaranteed high photosensitizer loading, while UCNP can upconvert NIR light into a light needed to activate these agents. These nanosensors showed marked selectivity and sensitivity for sensing the level of hypoxia in the cerebral anoxia model of zebrafish.

NIR-resonant nano-agents are widely employed in PTT in cancer. It depends on absorbing NIR and converting it to heat that eradicates the tumor cells [61, 62]. Novel NIR-induced DNA hybrid gated nanocarrier-based mesoporous coated with photothermal agent $\text{Cu}_{1.8}\text{S}$ nanoparticles were fabricated by Sun et al. [63]. The produced photothermal effect was directed towards DNA double strands and causes its denaturation. Furthermore, it triggered the doxorubicin/curcumin release that causes marked synergistic impact in the anticancer therapy nanoplatform. Yang et al. enhanced the cellular uptake and anticancer targeting therapy by synthesizing NIR triggered MoS_2 nanosheets coated MSNRs, modified with human serum albumin, and loaded with Ce6 [64]. In addition to NIR light-stimulated photodynamic therapy (PDT) and photothermal therapy (PTT), this carrier showed excellent biocompatibility, high contrast X-ray computed tomography imaging ability, in vivo fluorescence, and PDT synergistic effect from hyperthermia generation by MoS_2 .

Escoto et al. synthesized MSNs capped with gold nanoparticles for photo-triggered controlled drug release [65]. They found that the MSNs were efficiently linked with gold nanoparticles without the premature release of the paclitaxel into the normal healthy cells. A novel near-infrared laser-responsive mesoporous silica nanocarriers were composed of gold nanorods capped with reversible single-strand DNA. This DNA strand was controlled via a laser switcher that subsequently controls the release of drug from the nanocarrier. These nanocarriers demonstrated precise drug delivery at a specific time and to a particular region [54]. The isomerization of the photoactive sites controls the morphology of the photoactive molecules. Azobenzene is one of the most commonly used light-sensitive molecules due to its ability to

undergo reversible photoisomerization under UV irradiation [66]. A novel strategy for NIR induced anticancer drug release was introduced by Liu et al. [67]. They fabricated azobenzene-installed MSNs with up-converting nanoparticles structure (UCNPs). Owing to light converting capabilities of UCNPs and trans-cis photoisomerization of azo molecules, at 980 nm, the drug release was well-controlled and achieved better effective chemotherapy. Perylene-functionalized poly(dimethyl aminoethyl methacrylate) have been conjugated electrostatically to MSNs for controlled anticancer drug release. Perylene-functionalized poly(dimethyl aminoethyl methacrylate) were sensitive to visible light, and pH causing unsealing of the nanopores, and this enhanced the drug release because of synergistic effects [68].

3.1.4 Ultrasound-triggered drug release

Due to the fact of noninvasiveness and capability of penetration deeply in the tissue without damaging them, ultrasound is considered as one of the most promising triggers for drug release nanocarriers [69]. It has been reported that ultrasonic radiation improves the drug release from both biodegradable and non-biodegradable matrices. For instance, ultrasound triggered-MSNs were synthesized by grafting poly(2-(2-methoxy ethoxy) ethyl methacrylate-co-2-tetrahydropyranyl methacrylate) on its surface [70]. This copolymer consisted of both ultrasound and temperature triggered monomers. Upon ultrasound irradiation for 10 min, the hydrophobicity of the polymer changes leading to opening the pores of MSNs and releasing the drug in a sustained pattern. Similarly, at 4 °C, the conformation changes resulting in drug diffusion.

3.2 Endogenous stimuli-triggered drug release

Endogenous stimuli (biological stimuli) are the inherent environment of tumor tissues. For example, enzyme presence depends on acidic conditions and glucose concentration. In this section, we will discuss different responsive designs of nanocarriers to variations on pH value, enzyme, redox potential, and glucose.

3.2.1 pH-triggered drug release

The controlled drug released in the response of pH is one of the most exciting approaches. Drug release in the response of pH is a promising approach in cancer therapies because of the acidic microenvironment of the tumor cells, and pH change resulted from cancer cells internalization [71]. Generally, induced drug release from MSNs was accomplished by several strategies such as pH-cleavable linkers, acid-sensitive inorganic substances, polyelectrolyte, and supramolecular nanovalves. In 2009, pH-responsive mesoporous silica spheres

were synthesized by coating MSNs with poly (methacrylic acid-co-vinyl triethoxysilane) (PMV). The PMV shells manage the transfer of the drug molecules in response to pH-change [72]. Furthermore, pH-sensitive cleavable linkers were conjugated to MSN pores via hydrazine bonds for doxorubicin delivery (prodrug strategy). It was concluded that pH-directed drug delivery was highly effective with reduced doxorubicin systemic toxicity. Drug release in a sustained manner successfully reduced the unspecific secretion due to enzymatic hydrolysis. Moreover, this approach results in high drug loading, releasing efficiency, and cellular absorbance [73]. A polyacrylic acid grafted to MSNs was used to deliver and control the premature release of doxorubicin in bone cancer therapy [74]. Controlled pH triggered the release of drug molecules was achieved by acid-labile acetal linker and gold nanoparticles capped onto the MSNs. The release is based on acetal linkage cleavage at various pH values [75]. ZnO quantum dots were used to cap MSNs for effective intracellular dissolving in the acidic parts of cancer cells [76]. Recently, Nasrallah et al. designed core-shell magnetic mesoporous silica microspheres with a favorable pH-responsive porous shell for drug delivery [77].

3.2.2 Enzyme-triggered release

Enzyme-responsive MSNs have also been widely investigated in the past decade. Hydrolyzed starch attached on the surface of MSNs was introduced as a potential carrier to release the entrapped drug in the presence of the appropriate enzyme. Different delivery profiles were obtained by changing the degree of starch hydrolyzation (i.e., associated with the length of the saccharides molecules) [78]. Lactose-capped MSNs were first introduced by Amorous and co-workers in 2009 [79]. They are selectively uncapped using β -d-galactosidase by the dissociation of a glycosidic bond and hydrolysis of the disaccharide lactose into the monosaccharides galactose and glucose.

Cancer is known as overexpressing and down in regulating various enzymes; hence, MSNs can be used as release triggers for this concept. MSNs act via ester, peptide, and urea bond cleavage. Enzymes such as esterase, elastase protease, and matrix metalloproteinases (MMPs) are used. MMPs, especially MMP2 and MMP9, are employed in almost all tumor cells and accompanied by cancer metastasis and angiogenesis [80, 81]. A case in point, MSNs were anchored to avidin via MMP9 to target overexpressing MMP9 cancer sites specifically. Successful MMP-9-triggered drug release was achieved in patient tumor cells and mouse lung cancer [82]. Cathepsin B is a type of protease that has widely applied in different enzyme-triggered drug release systems. MSNs-dependent cathepsin B were synthesized by Cheng et al. [83].

3.2.3 Redox-triggered release

Redox triggered carrier is considered as one of the most efficient tumor therapeutic approaches. This approach is based on the significant difference in redox potential between cancer and healthy tissues. The amount of glutathione (GSH) in tumor cells is threefold higher than that in normal cells [84]. The existence of GSH biological reducers efficiently cleave disulfide bond (S-S), permitting the use of redox-triggered drug release systems. The first redox-triggered drug delivery system was reported by Lai et al. [85]. They used CdS nanoparticles to seal the pore access of MSNs via a disulfide bond. Several inorganic nanoparticles served as nano valves to trap the drug inside the MSNs via covalent bonding with disulfide linkers such as zinc oxide [86], iron oxide [87], and CdS [85]. In another study, MSNs were synthesized with disulfide redox cleavable linkers on the surface and conjugated to beta-cyclodextrin. The MSNs successfully controlled the drug release via this linker [88].

Redox responsive MSNs were also functionalized with polymers such as polyethylene glycol (PEG) using a disulfide linker. The drug release was significantly increased with an elevated amount of glutathione; on the other hand, the sealed PEG in reduced GSH acted as gatekeeper [89, 90]. Furthermore, several supra-molecules were used as gatekeepers on MSNs. For instance, adamantane was grafted onto the nanopores of drug-loaded MSNs using a disulfide linker. The efficient internalization of drug-loaded MSNs into the cytoplasm of tumor cells was accomplished. The high amount of GSH removed the adamantane caps by breaking disulfide linkers [91, 92].

3.2.4 Glucose-triggered release

Glucose-triggered release systems have received a lot of attention recently in drug delivery strategies [93]. Glucose-responsive MSNs are used in diabetes treatment by releasing a monitored amount of insulin in response to blood glucose concentration. For example, Zhao et al. synthesized boronic acid-functionalized MSNs for controlled delivery of both insulin and cyclic adenosine monophosphate (cAMP) in the response of glucose [93]. Gluconic acid-modified insulin was conjugated on the external surface of MSNs, which served as caps to trap the cAMP molecules inside the mesopores of MSNs. The presence of boronic acid resulted in the formation of more stable cyclic esters when exposed to glucose, leading to a controlled release of insulin and cAMP. Similarly, glucose-responsive alizarin functionalized MSNs integrated both drug and fluorescent real-time controlling issues [94]. Gluconate insulin using benzene-1,4-diboronic acid is integrated with the alizarin MSNs, which served as hypoglycemic drugs and gatekeepers. The glucose cleaved the boronate ester between alizarin and di-boronic acid resulting

in pores opening and fluorescence disappearance. Recently, intestinal permeation and oral delivery of insulin (10 U Kg^{-1}) have been significantly improved by incorporating anionic small silica nanoparticles ($< 200 \text{ nm}$) in both healthy and diabetic mice. The enhanced effect was nontoxic and reversible; in addition, the sustained hypoglycemic effect was achieved and lasted for a few hours [95].

3.2.5 Adenosine triphosphate (ATP)-triggered release

ATP is a multifunctional nucleotide responsible for providing energy for all biological processes. The ATP content in intracellular cytosols is higher than that in extracellular ones [96]. Therefore, ATP can be used to trigger the drug release inside the cytosol. Many reports confirmed the overexpression of tumor cells to ATP and the uncontrolled cancer growth [97, 98]. Graphene quantum dots were capped onto fluorescent MSNs through ATP aptamer for intracellular drug release [99]. The drug delivery was monitored via fluorescence resonance energy transfer, where under extracellular environment (low ATP concentration), the MSNs pores are closed, and the drug is trapped inside the MSNs. On the other hand, in the ATP-rich cytoplasm, the ATP aptamer changes its conformation resulting in shedding the graphene quantum dots from MSNs and releasing the drug in the tumor cells. Table 1 summarizes various stimuli systems employed for drug delivery using MSNs.

4 Targeting approaches

4.1 Passive targeting

The mechanism of passive targeting was discussed for the first time by Mastumura in 1986. It is mainly based on the leaky discontinuous microvasculature and defect of lymphatic functions of tumor tissues that result in improving permeability and retention (EPR) effect. Tumors exhibit rapid angiogenesis to maintain its proliferation by delivering the needed amount of oxygen and supplies [100]. This rapid angiogenesis results in apertures that enable the synthesized nanocarriers to enter the tumor cells and accumulate in it. However, this accumulation would take place if MSNs can escape from fast renal clearance and reticuloendothelial systems. This approach has been widely used to improve targeting using silica nanocarriers [101]. To illustrate, MSNs were synthesized by integrating functional nanoparticles in mesoporous shells included magnetic nanoparticles, and then attaching it onto the surface of MSNs. These MSNs successfully accessed the tumor sites of mice via passive targeting. In addition, these MSNs are applied in fabricating multimodal imaging probers for MR [102]. In the next sections, the different factors control

the tissue accumulation, and cellular uptake of MSNs will be introduced.

4.1.1 Size and shape

Nanoparticle size determines circulation, distribution of MSNs, tissue accumulation, and cellular uptake. The agreed optimum nanoparticle size for biomedical applications is ranged from 10 to 300 nm. However, maximum internalization was achieved from nanoparticles of 100–200 nm size by obtaining maximum accumulation in tumors via EPR [103, 104]. Small particles of 80 nm size have the longest mean residence time (2 h) and slow biodegradation. Nanoparticles larger than 400 nm display the inability to penetrate the tumor tissues and offering therapeutic effects. Spherical MSNs of 80–360 nm size are mainly distributed in the liver and spleen [105]. Lou et al. studied the cellular uptake of MSNs in HeLa cells [106]. The cellular uptake was found to be size-dependent, and particle size of 50 nm is optimum to guarantee efficient cell internalization. In comparison, particles tend to agglomerate when particle size is less than 50 nm, causing endocytosis lowering [107].

In addition, the shape of the nanoparticles represents an essential role in EPR-dependent tumor passive targeting. Gao et al. proposed the shape impact on cell internalization [108]. It was hypothesized that upon membrane stimulation, a diffusive ambulant receptor wraps the ligand coated spherical particles. Generally, spherical nanoparticles have been widely used in many biomedical applications because of its simple fabrication method and shorter wrapping time that resulted in faster internalization [108]. The shape influence of silica nanoparticles on cell internalization efficiency was discussed for the first time by Trewyn et al. [109]. They designed spherical and tube-like MSNs labeled with fluorescein and tested them in cancer cell lines and noncancerous. The results demonstrated that cellular uptake and endocytosis are morphology dependent. Similarly, spherical MSNs demonstrated faster internalization rates in Chinese hamster ovarian than tube-like ones [109]. Aspect ratios (ARs) showed a positive association with endocytosis rates and accumulation of MSNs in the cell [110]. This observation was also confirmed by designing three MSNs with different aspect ratios (1, 2, and 4) while maintaining the diameter, chemical structure, and surface charge unchanged. These MSNs offered different internalization efficiency in A375 human melanoma. However, particles with larger ARs provided higher cellular uptake [110]. Another report demonstrated that rod-like MSNs with ARs of 2.1–2.5 displayed faster internalization and more efficient penetration than the spherical ones [111]. Short rod-like MSNs existed mainly in the liver with a rapid clearance rate while the long ones distributed in the spleen [112]. Based on these research findings, it is hard to conclude the most suitable MSNs shape needed for passive targeting systems.

Table 1 Reviewed stimuli employed for drug delivery using MSNs

Stimuli	Mechanisms	Reference
1. External stimuli		
Temperature	MSNs coating with PNIPAM	[38]
	Dual responsive MSNs (Magnetic field + temperature)	[39]
Magnetic field	Increase of temperature under the action of an alternating magnetic field	[40–42, 51]
	Superparamagnetic iron core in MSNs	[46, 48, 49]
	Magnetic MSNs+ cyclic amino acid lyp	[50]
Light	Magnetic MSNs + quantum dots as a cap	[54, 61–64, 67]
	NIR	[59, 65]
	MSNs capped with gold nanoparticles	[56, 58, 60]
	Activated via photosensitizer agent	[68]
Ultrasound	Dual responsive MSNs (NIR + pH) Perylene-functionalized poly (dimethyl-aminoethyl methacrylates)	[70]
	-MSNs were synthesized by grafting 2-(2methoxyethoxy) ethyl methacrylate co-tetrahydro-pyranyl methacrylate on its surface	[70]
2. Internal stimuli		
pH	PH-responsive mesoporous silica spheres, accomplished by coating with poly (methacrylic acid-co-vinyl triethoxysilane) (PMV).	[72]
	Conjugating pH-sensitive cleavable linker to MSN pores via hydrazine.	[73]
	A poly acrylic acid grafted to MSNs	[74]
	Acid- liable acetal linker and gold nanoparticles capped onto the MSN.	[75]
	MSNs capped with ZnO (Acidic cleavage)	[76]
Enzyme	Hydrolyzed starch attached to MSNs surface	[78]
	Lactose capped MSNs	[79]
	MSNs anchored to avidin via MMP9, specifically target over expressing-MMP9 cancer sites	[82]
Redox	MSN-dependent cathepsin B	[83]
	S-S cleavage disassembly	[89–92]
Small molecules	Glucose concentration	[93, 94]
	Anionic small silica nanoparticles (oral insulin delivery)	[95]
	Competitive binding in presence of ATP	[99]

ATP, denosine triphosphate; *pNIPAM*, poly(N-isopropylacrylamide); *NIR*, near infrared radiation; *ZnO*, zinc oxide

4.1.2 Surface

MSN surface plays a vital role in cell internalization, biodistribution, cell binding, and blood circulation time. MSNs are well-known to acquire large surface area and pore volume. The challenges are faced in designing MSNs with appropriate functionalization and charges to ensure the proper interaction with the targeted physiological cells [113]. Interaction between red blood cells (RBCs) and silica nanoparticles resulted in hemolysis. The negative charge of the silanol group results in strong affinity and binding to RBCs. These underlying phenomena affect the internalization of MSNs in the tumor tissues and compromise the efficiency of passive targeting. However, this behavior could be modified by attaching an organic constituent to MSNs [114, 115]. Additionally, the resulted opsonization from nonspecific adsorptions of plasma proteins on the MSNs surface affects the

passive targeting. This adsorption creates protein corona that rapidly cleared via RES system [116]. PEGylation is a common strategy used to overcome these phenomena. PEGylation can be achieved by surface functionalization of MSNs using polyethylene glycols (PEGs) [117]. This functionalization process reduces the protein binding via electrostatic repulsion and steric hindrance effects. The optimum PEG molecular weight for this purpose is 10,000–20,000 g/mol. Moreover, PEGylated MSNs displayed a significant reduction in hemolysis events [118]. Recently, the influence of surface modification of MSNs using lipids and PEGs on mesenchymal stem cell internalization was investigated [119]. This modification resulted in improving internalization capacity 17 times higher (> 80%) than the unmodified ones. On the other hand, the protein resistance character was reduced at temperatures higher than 35 °C. Moreover, the pharmacokinetics property of PEGylated MSNs was altered over time with repeated

injections in animals [120]. It was emphasized that PEGs produce anti-PEG IgM that resulted in faster blood elimination [120]. Recently, the conjugation of MSNs with zwitterionic copolymers resulted in enhancing the stability and increasing blood circulation half time [121]. Lipid coating has also been applied as an alternative strategy to escape from RES clearance [122]. The charge on the surface of the nanoparticles plays a significant role in passive targeting. Since the cell membranes are negatively charged, positively charged surface nanocarriers offer higher cell uptake than the negatively or neutrally charged ones. The final surface charge of nanocarriers tends to be negative due to plasma coating [123]. Surface functionalization of MSNs with contrast agent indocyanine green (ICG) was investigated for photoacoustic imaging to overcome ICG serum protein binding problems and rapid clearance [124]. NH_2 modification of MSNs displayed higher ICG loading ability (16.5 wt%) compared to PO_3 modified MSNs (3.5 wt%). Furthermore, NH_2 MSNs showed 1.34-fold PA signal improvement compared to ICG alone. MSNs modification with silanes having NH_2 or PO_3 groups resulted in improving the solubility of Vorinostat for the treatment of cutaneous T cell lymphoma (3.9-fold and 4.3-fold respectively, compared to the free drug) [125]. Furthermore, resveratrol was efficiently encapsulated with NH_2 and PO_3 modified MSNs. These MSNs showed dose-dependent sensitization of docetaxel for the hypoxic condition in prostate cancer [126].

4.2 Active targeting

Active targeting relies on the interaction between cancer cells and drug loading cargos. Functionalizing the surface of MSNs with a specific ligand is necessary for the active targeting of cancer cells. Macromolecules, antibodies, nucleic acids, vitamins, sugars, and peptides are examples of ligands that applied to anchor the MSNs for successful active targeting. This strategy depends on the ligand affinity to the cancer cells overexpressed receptors [127]. In general, the site-specific active targeting strategies can be classified into a tumor cell, vascular, and subcellular organelles targeting (Fig. 7).

4.2.1 Surface functionalization of MSNs for cancer applications

4.2.2 Functionalization by folic acid

Targeting using folic acid has been widely used in many *in vitro* studies, yet there are limited reports discussed in *in vivo* studies. To illustrate, docetaxel-loaded folic acid MSNs showed improved cellular uptake, apoptotic, and anti-cancer activity against breast cancer MCF-7 cells. The *in vivo* fluorescent imaging revealed the accumulation of folic acid MSNs in the kidney of healthy mice [129]. Other conducted

studies compared the performance of camptothecin-loaded MSNs to folic acid functionalized camptothecin/MSNs in terms of xenograft pancreatic cancer suppression. Folic acid-functionalized camptothecin/MSNs provided excellent biocompatibility and significant inhibition of tumor growth [130]. Naturally originated anticancer prodrug colchicine, quercetin, and curcumin have been loaded on folic acid conjugated MSNs. These novel MSNs were successfully employed in targeting *in vitro* hepatocellular cancer cells (HeLa). Additionally, they showed marked enhanced targeting to liver cancer cells [131].

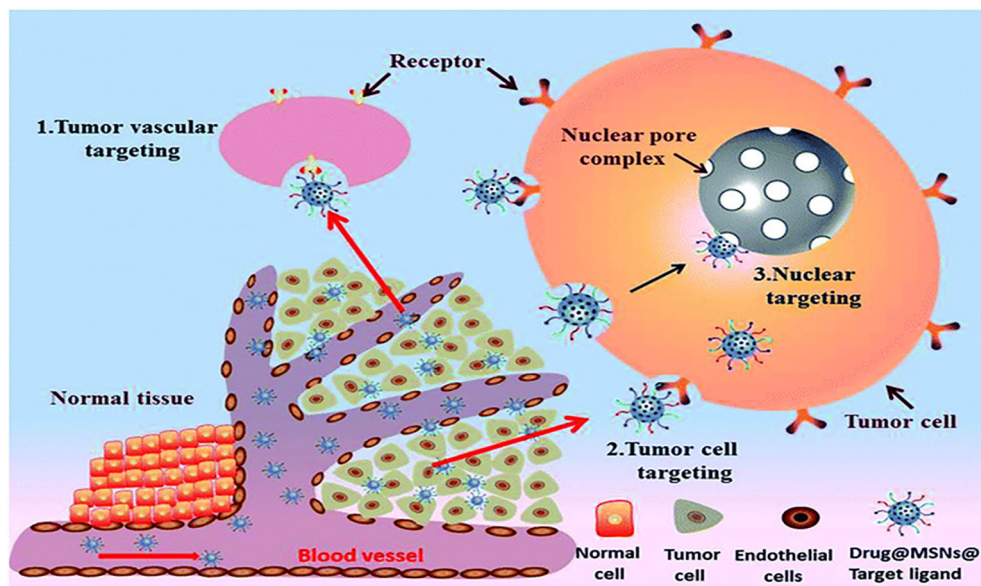
4.2.3 Functionalization by mannose

Another promising approach is the functionalization of MSNs using mannose. Anchoring mannose to MSNs with photosensitizer showed remarkable targeting breast cancer cells (MDA-MB-231). Mannose exhibited high photodynamic efficiency on breast cancer cells [132]. Cancer cells were demonstrated to have a high affinity to glycosylated moieties. The galactose-conjugated fluorescent nanoparticles were designed using lactobionic acid (LA) through EDAC linkage. They are highly selective to liver cancer cells without harming the normal ones [133]. Besides, colorectal cancer cells can be targeted using galactose-functionalized camptothecin/porphyrin-loaded MSNs. This synergistic combination of photodynamic therapy (porphyrin) and targeted drug delivery (camptothecin) enhanced the internalization of MSNs in the endosome [134].

4.2.4 Functionalization by monoclonal antibody

The monoclonal antibody can target cell receptor antigens that are overexpressed on cancer cells more than that overexpressed on the normal ones. For this purpose, a monoclonal antibody (mAb) functionalized MSNs containing green fluorescent dye were prepared by grafting Herceptin to MSNs outer surface for selective targeting epidermal growth factor receptor HER2/neu glycoprotein. The synthesized nanocomposites showed high internalization in breast cancer cells as well as selective targeting by Herceptin [135]. Epithelial cell adhesion molecule antibody (EpCAM) is a transmembrane glycoprotein, which is overexpressed in many cancer cells and cancer stem cells. EpCAM acts as an efficient preventive of metastatic cancer by targeting the circulating tumor cell and inhibiting the contact-invasion to vascular intima [136]. Interestingly, long-circulating (EpCAM) grafted to mifepristone (MIF)-loaded MSNs prevented lung metastasis cascade in the mouse. This system selectively targets colorectal cancer cells in both cell and blood mediums [137]. Similarly, doxorubicin-loaded MSNs anchored to EpCAM-aptamer were efficiently delivered to colon EpCAM overexpressing SW620 tumor cells. This unique design exhibit a controlled

Fig. 7 Schematic diagram illustrating active targeting by MSNs. (1) Cancer vascular targeting. (2) Cancer cell targeting. (3) Nuclear targeting (reprinted with permission from [128])



release pattern, and the drug release was favored in acidic medium (pH 5.5) resulted in improving targeted therapy and reducing side effects [138]. Mucin-1 is also a transmembrane glycoprotein that is overexpressed nearly in all epithelial cancer cells. Near-infrared MSNs were linked to the mucin-1 antibody successfully to detect primary breast carcinoma and to track breast cancer cell metastases in transgenic mouse model [139]. Recently, targeting cancer stem cells have grabbed researchers' interest in initiating and maintaining tumor proliferation. For instance, daunorubicin-loaded MSNs were conjugated to succinic anhydride capped with the anti-B220 antibody for targeting leukemia stem cells. Also, incorporating these MSNs markedly retarded leukemia proliferation in mice model [140].

4.2.5 Functionalization by DNA aptamer

Recently, the functionalization using signal responsive DNA switches has gained a lot of attention, making them valuable alternatives to antibodies and peptides. Various triggers can be used to stimulate the DNA switches such as light, metal ions, pH, and aptamer substrate complexes [141]. The release of the substrate entrapped in MSNs is controlled by tailoring nucleic acid tags. In normal cases, this tag inhibits the release of the substrate from the MSN pores. On the other hand, the presence of a biomarker induces a catalytic decomposition resulting in nucleic acid rearrangement and pore opening, permitting the substrate release [142]. Nucleic acid cap identifies molecular biomarkers via aptamer substrate complexes [142]. Mucin-1 is overexpressed in about 90% of breast cancer therapies. Safranin O-loaded MSNs have been capped with negatively charged aptamers for efficient internalization in MDA-MB-231 breast cancer cells and controlled substrate release.

Furthermore, the MSNs were radiolabeled with ^{99m}Tc for improved carcinoma targeting [143].

4.2.6 Functionalization by hyaluronic acid

Hyaluronic acid (HA) is the main component of the extracellular matrix of the biocompatible glycosaminoglycan needed for growth and maintaining tissue stability. Many tumor cells, such as CD44, have been reported to have overexpression to HA. Besides, HA exhibits multifunctional moieties available for binding several anticancer drugs and nanocarriers. Thus, HA functionalized MSNs are widely used in various pharmaceutical applications [144]. Colon cancer cell (HCT-116) surface exhibits CD-44 overexpression. Therefore, it has been targeted using HA-conjugated doxorubicin-loaded MSNs. High apoptosis to HCT-116 and enhanced internalization in the tumor cells was achieved [145]. Lee et al. investigated for the first time the antiviral characteristics of functionalized MSNs for herpes simplex virus infection [146]. They demonstrated that glycosaminoglycans mimetic functionalized MSNs act as viral binding/entry inhibitors that prevent the cells from being infected by HSV-1 and HSV-2.

4.2.7 Functionalization by peptides

The integrin receptor is overexpressed in various cancer cells, and this receptor could be selectively targeted using tripeptides (arginine-glycine-aspartic acid) (RGD). For instance, MSNs were designed for NIR-responsive drug delivery and chemo/photothermal therapy. RGD sequences were anchored on the doxorubicin-loaded MSNs surface, and they showed efficient internalization in tumor cells through receptor-mediated uptake and high stability in the bloodstream [147]. RGD anchored to MS-encapsulated gold

nanorods offered an improved uptake in breast carcinoma via alpha V beta3 integrin, and they significantly enhanced the radio-sensitization of triple-negative breast cancer cells [148]. ADDA-TCPP amphiphile peptides were immobilized onto doxorubicin-loaded MSNs. These MSNs blocked the premature release of doxorubicin via the hydrophobic interactions between alkyl sequences of ADDA-TCPP. Additionally, the release of the drug was triggered by glutathione [149]. Moreover, MSNs showed promising potential for guided tissue regeneration [150]. Xu et al. fabricated MSNs by coaxial electrospinning, where both growth and antibiotic factors were loaded in the core and shell of MSNs, respectively. These MSNs demonstrated sustained release patterns of the growth factor besides efficient ontogenetic regeneration abilities towards bone marrow stromal cells (BMSCs) as well as excellent antibacterial properties.

4.3 Vascular targeting

Angiogenesis (formation of new blood vessels) is necessary for the growth and metastasis of cancer cells [151]. Therefore, vascular targeting is considered as a promising alternative for targeting solid tumors. This strategy is based on indirect targeting carcinomas through destructing endothelial cells of tumor vessels to cease their needed oxygen and nutrients [152]. Vascular targeting is more favored than other approaches as cellular internalization in tumors and extravasation from blood vessels are not required [153]. Therefore, many challenges can be overcome using this strategy, such as tumor heterogeneity and poor delivery. Although the concept of vascular targeting was introduced in the 1920s [154], but the first successful *vivo* cancer vascular targeting was achieved by Burrows and Thorpe in 1993 [155]. This approach is based on conjugating the nanocarriers with vasculature-targeting ligands such as vascular endothelial growth factor (VEGF) and arginine glycine aspartic acid (RGD) peptides. In addition, VEGFR-targeted MSNs anchored to Cu-NOTA/PEG-VEGF were synthesized [154]. These MSNs displayed threefold improvement in terms of cancer accumulation [156]. Similarly, conjugating ursolic acid and VEGF-targeted siRNA to MSNs anchored with folic acid were examined on folate receptor overexpressing HeLa cells and folate receptor-negative HepG2. They combined active and passive targeting concepts and achieved a significant improvement in targeting efficiency. Furthermore, the stability of VEGF-targeted siRNA was enhanced using folic acid ligand [157]. In summary, incorporating suitable ligands results in improving targeting efficiency and overcoming the existing challenges.

4.4 Nuclear targeting

The concept of nuclear targeting using MSNs has been introduced by Pan et al. [158]. MSNs were conjugated to nuclear

localization signal TAT peptides. The MSNs-TAT systems of 50-nm diameter were able to target the nucleus and accumulate doxorubicin in the tumor nucleus. This approach has been developed to overwhelm many challenges. For instance, several free anticancer drugs provided a suppressed therapeutic efficiency after reaching the nucleus owing to many physiological barriers [159]. The nuclear-targeted strategy is adhered via anchoring MSNs surface to nuclear localization signals such as gold nanoparticles [160, 161], magnetic nanoparticles [162], and quantum dots [163]. Recently, this approach has been widely applied in gene therapy [164]. To illustrate, functionalized MSNs were employed in codelivery of osteogenic genes and drugs. This dual system showed synergistic improvement in the osteogenic impact of stem cells. Moreover, doxorubicin delivery via nuclear targeted MSNs overcome the multidrug-resistant problem (MDR) by upregulating the genes responsible for the apoptosis of multidrug-resistant cells. Additionally, this approach accomplished downregulated expression of the apoptosis-suppressing gene *bcl-2* and prevented the activation of the DNS repair system for improved MDR cells apoptosis [165].

4.5 Multistage targeting

This approach was recently developed due to unsatisfactory therapeutic outcomes resulted from single targeting ones (cancer cells, nuclei, or endothelial cells). This novel strategy can be achieved via conjugating multiple ligands on the surface of MSNs exhibiting high selectivity and affinity to different tumor types. This design results in targeting various disease sites, a high therapeutic index, and reduced systemic side effects [166]. Multistage ongoing targeting approach relies on synthesizing magnetic MSNs for efficient accumulation in the cancer tissues, then conjugating TAT peptide to MSNs surface for nuclear targeting, and finally attaching charge-conversional polymer originated from folate anchored chitosan. This smart approach causes accumulation of MSNs in the tumor using magnetic nanoparticles, enhances cellular uptake via folate receptor endocytosis, and targets cell nucleus using TAT peptides [167]. Pan et al. have extrapolated the multistage targeting approach to *vivo* application. MSNs were coupled with RGD peptide that resulted in efficiently targeting the tumor vasculature and cell membrane [168]. Additionally, TAT peptides were used for nuclear targeting signals that guaranteed efficient nuclear uptake. Significant enhancement in therapeutic efficacy and tumor suppression (98.6%) were noticed as a result of sequential vascular-to-cell nuclear-targeted drug delivery. Naz et al. developed novel multistage-targeted anticancer MSNs that are capable of targeting both CD44 and mitochondria [169]. Utilizing triphenylphosphine (TPP) as a mitochondrial targeting agent was anchored onto the surface of MSNs. Doxorubicin was loaded in MSNs, and HA was capped with MSNs (as a

tumor-targeting agent). CD44 receptor of tumor cells uptake these MSNs composite, then accumulated in mitochondria due to the presence of the TPP compound. HAase enzyme overexpression in cancer cells caused degradation of HA and release of doxorubicin and resulted in efficient eradication of cancer cells. Hollow MSNs anchored with tLyp-1 peptide (tHMSN) for dual-targeting drug delivery to cancer cells, and angiogenic blood vessels were synthesized by Liu et al. [170]. Doxorubicin loaded on tHMSN showed a high inhibitory effect on MDA-MB-231 cells and HUVECs in vitro. In another study, cell-targeting agents, folic acid, dexamethasone, and nucleus targeting agent (TAT) were coupled in 40 nm MSNs [171]. These cell-nucleus sequential targeting-based MSNs improved the inhibition efficacy of doxorubicin on Hela cells and decreased the toxic side effects on normal cells.

5 Conclusion

It is evident from ineffective conventional therapies that there is a desperate need for distinct and promising therapies. As extensively discussed, MSN-based nanostructure technology has the potential to play critical functions in cancer treatment. MSNs have promising potential in drug delivery due to its unique physical-chemical characteristics. Through functionalization with various groups and molecules on MSNs surface, stimuli-responsive and targeting approaches have been proved to provide unprecedented opportunities and advances for novel theranostic modalities. However, despite the extraordinary breakthrough that has been made to modify and enhance MSNs based drug loading and release, still, using MSNs in clinical applications require more efforts to translate these preclinical findings into the clinic. Currently, the most significant studies are achieved as in vitro experiments (cell culture), while tissue penetration is not demonstrated yet. Such results may not mimic the in vivo environment and described critical difficulties in translating these MSNs based nano-systems in the clinical trials. Furthermore, it is widely noticed that very few articles discussed the fate and the clearance pathways of these nanomaterials that may be accumulated and caused massive toxicity. Therefore, more preclinical in vivo animal studies need to be conducted to demonstrate various unknown systemic aspects of these nanocarriers, such as processing, accumulation, and elimination in the living organisms.

Nanostructured materials, namely mesoporous silica nanoparticles, represent a promising candidate in drug delivery applications. In this review, we highlighted the recent insights in the functionalization of MSNs for cancer therapy applications, including stimuli triggered drug release and cancer-targeted delivery. We also reviewed the current designs applied for successful targeting using MSNs. Prospective

investigations are still needed to ensure the introduction of MSN-based drug delivery systems to the clinical market.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

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