Active scaffolds for on-demand drug and cell delivery

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Abstract

Porous biomaterials have been widely used as scaffolds in tissue engineering and cell-based therapies. The release of biological agents from conventional porous scaffolds is typically governed by molecular diffusion, material degradation, and cell migration, which do not allow for dynamic external regulation. We present a new active porous scaffold that can be remotely controlled by a magnetic field to deliver various biological agents on demand. The active porous scaffold, in the form of a macroporous ferrogel, gives a large deformation and volume change of over 70% under a moderate magnetic field. The deformation and volume variation allows a new mechanism to trigger and enhance the release of various drugs including mitoxantrone, plasmid DNA, and a chemokine from the scaffold. The porous scaffold can also act as a depot of various cells, whose release can be controlled by external magnetic fields.

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Alginate gels were covalently cross-linked with adipic acid dihydrazide (AAD) (Fig. 1D) (40), because the covalent cross-links allowed gels to maintain the macroporous structure following lyophilization and subsequent rehydration. Peptides containing the RGD amino acid sequence were covalently coupled to the polymer prior to gel formation (Fig. 1D) (44). The RGD coupling confers a specific mechanism for integrin-mediated cell adhesion to the otherwise nonadhesive polymer, and the RGD density can be manipulated to provide control over cell adhesion.

The influence of the macropores on the gel mechanical properties were next evaluated because the stiffness of the gels will dictate the extent of deformation that will result from a specific magnetic field. It is well established that by creating pores in a gel, one can greatly reduce the gel’s elastic rigidity (45). A ferrogel fabricated with 13 wt % Fe₃O₄ and 1 wt % alginate cross-linked by 5 mM AAD gives an initial Young’s modulus (i.e., the slope of the initial part of the stress vs. strain curve in Fig. 2A) of ∼30 kPa in a compression test. Introducing connected pores of ~700-μm diameter into the ferrogel led to a dramatic reduction in the initial modulus, to ∼2.5 kPa (Fig. 2A). These macroporous gels can also be reversibly deformed to a compressive strain of over 80% before fracture, in contrast to the more brittle nature of the standard nanoporous alginate gels (Fig. 2A). In addition, it has been observed that the initial Young’s modulus of the macroporous gel decreases with the rise of the freezing temperature used in preparation of the gel (Fig. S2). Macroporous ferrogels fabricated with 13 wt % Fe₃O₄ and 1 wt % alginate cross-linked by 5 mM AAD, and frozen at −20°C were used for the remaining experiments. The macroporous ferrogels with interconnected pores, larger pore size (Fig. 1B), higher concentration of iron oxide particles (Fig. 1C), and lower modulus (Fig. S2) were chosen because (i) drug and cell transport will presumably be more efficient in scaffolds with larger pore sizes and better connectivity (40), (ii) a higher iron-oxide-particle density gives a higher body force under the same magnetic field, and (iii) a gel with lower modulus tends to deform more when subject to the same body forces. Large deformation of the gel leads to more pronounced effects on drug and cell release.

The deformation of nanoporous and macroporous ferrogels under the influence of a moderate magnetic field was next examined. Subject to a nonuniform magnetic field, a ferrogel experiences a body force proportional to the gradient of the applied field, which results in a shape change (46, 47). A cylinder of a nanoporous ferrogel reduced its height by ∼5% when subjected to a vertical magnetic-field gradient of ∼38 A/m². The corresponding macroporous ferrogel deformed ∼70% under the same magnetic field. SEM images of a freeze-dried macroporous ferrogel in the undeformed and deformed states. (Scale bar: 500 μm.)
larger deformation, ∼70%, under the same magnetic field (Fig. 2C), due to its lower modulus and the increase of iron-oxide-particle density during deformation (Fig. 2A). In addition, the cylinder of the macroporous gel also reduced its volume by ∼70% (Fig. 2C), as contrasted to a minimal volume change of the nanoporous gels (Fig. 2B). The large volume change of the macroporous gels was caused by collapse of the pores in these gels, as demonstrated by SEM images of freeze-dried macroporous ferrogels in the deformed and undeformed states (Fig. 2D). The collapsing pores will force water contained in the connected pores to flow out of the gel. Once the magnetic field is off, the elastically deformed gel quickly returns to its original, undeformed configuration in less than 1 s, as surrounding water was reabsorbed into the gel (Movie S4).

We hypothesized that the magnetic-field-induced gel deformation and resulting water convection and associated shear forces could accelerate the release of drugs encapsulated in ferrogels or cells adherent to the pore surfaces of the macroporous ferrogels. To test this hypothesis, we first chose mitoxantrone (M, 444 444), an antineoplastic agent, as a model drug. Mitoxantrone forms an ionic complex with the carboxylate groups on alginate, which retards its release (48). Macroporous ferrogels each containing 300 mg mitoxantrone were placed in a PBS solution. Every 30 min, one group of the macroporous ferrogels was subjected to 120 cycles (on/off) of the magnetic field over 2 min by manually approaching and retracting a magnet against the gels, while no magnetic field was applied to the other gels. The short recovery time of the gels allows them to recover their initial dimensions in each cycle. Gel deformation and water convection promotes dissociation of drugs from the polymer and enhances transport of unbound drugs out of the gel. The macroporous gel deformed reversibly and promptly in response to the cycling magnetic field, and the applied magnetic stimulation greatly accelerated the release of mitoxantrone, as shown grossly by the increasing blue color in the PBS. (see Movie S2). The cumulative release profile showed a stepwise increment with magnetic stimulation, and the total amount of mitoxantrone released from the stimulated gel was ∼7 times more than that of the control case after 3 h (Fig. 3A).

This study was repeated with higher molecular-weight biological agents, plasmid DNA (M, 10⁸) condensed with polycyanoa- ronitrile (PEI, M, 22,000), and the chemokine SDF-1α (M, 8,000) to determine if this approach would have utility with a wide range of potential drugs. Again, significant stepwise increases in release were noted with application of the magnetic field, as compared to control gels (Fig. 3B and C).

Next, we used the macroporous ferrogel to examine controlled cell release in vitro (Fig. 4A). The RGD on alginate in macroporous ferrogels enables cells to adhere on the porous surfaces and the adhesion strength increases with RGD density (49). A baseline RGD density was set to 7.43 μmol RGD per gram of alginate (50), and 50% and 10% of the baseline RGD density were also used to examine the role of gel adhesiveness on cell release. Human dermal fibroblasts (1.5×10⁶) were seeded in each gel, and gels were incubated at 37°C for 4 h to allow cell adhesion. Thereafter, the macroporous ferrogels were subject to 120 cycles (on/off) of the magnetic field for ∼2 min, at 2-h intervals. The cumulative release of cells was quantified as a function of time (Fig. 4B). Magnetic stimulation resulted in burst release of cells and the release profile varied for gels with different RGD densities. After 6 h (three rounds of magnetic stimulation), a third of the cells were released from gels with the baseline RGD density, while over half of the cells were released from ferrogels with 50% less RGD. The gels with only 10% of the baseline RGD content delivered over 90% of their cells after 4 h (two rounds of stimulation). These results are consistent with the lower RGD densities providing weaker cell adhesion, resulting in more cell detachment and release under the magnetic stimulation (51). The released cells were collected and probed for viability. More than 95% of the released cells were viable (Fig. S3). The released cells were also incubated in a Petri dish at 37°C. These cells spread normally and became confluent after 2 d (Fig. 4C), indicating that released cells remained viable and functional.

Finally, we examined the capability of macroporous ferrogels in controlled delivery of cells in vivo. Mouse mesenchymal stem cells (1.5×10⁶) stained with DIOC18, a membrane dye with near infrared emission maximum, were seeded in macroporous ferrogels with 50% baseline RGD content. The gels were implanted into the subcutaneous space of mice. One hour after the implantation, the gels were subjected to 120 cycles (on/off) of the external magnetic field by approaching and retracting a magnet against the mouse skin over the gel. As a control, implanted ferrogels were subject to no magnetic stimulation, and in vivo fluorescence images of both conditions were obtained before and after magnetic stimulation (Fig. 5). The control mouse (Fig. 5A and B, Left) showed almost no change in fluorescence, indicating few cells were released from the ferrogel. In contrast, application of the external magnetic field led to a significant increase in

![Graph A](image1.png)

![Graph B](image2.png)

![Graph C](image3.png)

**Fig. 3.** (A) Cumulative release profiles of mitoxantrone from macroporous ferrogels subject to 2 min of magnetic stimulation every 30 min, or no magnetic stimulation. (B) Cumulative release profiles of plasmid DNA from macroporous ferrogels subject to 2 min of magnetic stimulation every 2 h or no magnetic stimulation. (C) Cumulative release profiles of SDF-1α from macroporous ferrogels subject to 2 min of magnetic stimulation every 2 h or no magnetic stimulation. Values represent mean and standard deviation of each increment (n = 3–5).
circles on the figure. The positions of the gel disks are indicated by.

The results of this study suggest an on-demand and reversible approach for delivering various biological agents from a newly designed macroporous ferrogel, in response to external magnetic fields. A delivery mechanism, based on deformation of macroporous ferrogels under magnetic stimulation, and subsequent water flow through connected pores is demonstrated. By creating large pores in a ferrogel, one can greatly reduce the rigidity of the gel and increase its hydraulic conductivity (45). As a result, the macroporous ferrogels demonstrate very large deformation, volumetric change, and water convection under applied magnetic fields, in contrast to the minimal values of conventional nanoporous ferrogels. Furthermore, it is generally difficult for nanoporous ferrogels to release drugs with high molecular weight, such as proteins and plasmid DNA, due to the limited mobility of these drugs in the nanopores (32–37). The connected macropores in the new ferrogel enable the rapid transport of various drugs ranging from small molecules such as mitoxantrone to very large molecules such as the protein SDF-1α and plasmid DNA out of the gel, upon magnetic stimulation. This result is consistent with previous observations that mechanical compression of macroporous or nanoporous gels can enhance the release of biological molecules (38–52).

Discussion

The study has been focused on the capability of active porous scaffolds to act as allow a mechanism for on-demand and cell release under the control of external magnetic fields. These scaffolds also act as active depots of various cells whose release can be controlled over time. The polymers used to form conventional ferrogels typically do not support cell adhesion (33, 37). To address this issue, the polymer used to form the ferrogels in this study was covalently coupled with cell-binding peptides because the peptide density can be manipulated to provide control over cell adhesion. On-demand release of human dermal fibroblasts and mouse mesenchymal stem cells were demonstrated in vitro and in vivo under the control of external magnetic fields. The peptide-modified macroporous ferrogel also maintains the adhesion and viability of the resident cells, and subjecting the gels to external magnetic stimulation allows one to release a prescribed number of the resident cells on demand over various time frames. Multiple parameters in this delivery system are tunable, including peptide density, the strength of applied magnetic field, number of magnetic cycles, and frequency of magnetic stimulation, to control the release of various cell types. This on-demand release of cells from porous scaffolds is potentially of wide utility for tissue regeneration and cell therapies. The current study has been focused on the capability of active porous scaffolds in on-demand drug and cell deliveries, but we also expect these macroporous ferrogels to find broader applications, including serving as actuators and sensors in biomedical and other applications, due to their large and fast deformation and volume change under magnetic fields.

Materials and Methods

Materials. Sodium alginate with high guluronate content (Protanal LF 20/60) was purchased from Pronova Biopolymers, Inc. and used without further purification. PBS was purchased from Invitrogen. AAD, 1-ethyl-3-(dimethylamino)propyl) carbodiimide (EDC), MES, 1-hydroxybenzotriazole (HOBt), Iron(III) chloride hexahydrate, oleic acid, 1-octadecene, and cetyltrimethylammonium chloride were purchased from Sigma-Aldrich. Sodium oleate was purchased from TCI. Pluronic F127 was purchased from BASF. The integrin-binding peptide (Gly)₄-Arg-Gly-Asp-Ala-Ser-Ser-Lys-Tyr was purchased from Commonwealth Biotech.

Macroporous Ferrogel Fabrication. The Fe₃O₄ nanoparticles were fabricated according to ref. 42 and the surfactant, Pluronic F127, was coated on the nanoparticles following the procedure in ref. 43. Alginites were coupled with RGD following aqueous carbodiimide chemistry as described in ref. 44. To prepare macroporous ferrogels, RGD-modified alginate in MES buffer (0.1 M MES and 0.5 M NaCl, pH 6.0) was sequentially mixed with an aqueous solution of Fe₃O₄ nanoparticles, HOBt, EDC, and AAD. The concentration of alginate was 1 wt %, Fe₃O₄ nanoparticles was 13 wt %, and AAD was 5 mM in the resulting solution. A baseline RGD density was set to 7.43 μmol RGD per gram of alginate according to ref. 50. The mixture was immediately cast between two glass plates separated by 4–11 mm.

Fig. 4. (A) Schematic plot of gel deformation and resulting water convection inducing cell release from macroporous gels (51). (B) Cumulative release profiles of fibroblasts from macroporous ferrogels with 100% (cross), 50% (circle), and 10% (square) of the baseline RGD density, following application of cycled magnetic field. (C) Spreading and proliferation of the released cells at various times after replating onto tissue culture plastic. Values in B represent mean and standard deviation of each increment (n = 3–5). Differences between the values of cell release at the various RGD densities were statistically significant at each time point (p < 0.05).

Fig. 5. In vivo fluorescence images of mice implanted with macroporous ferrogels containing mouse mesenchymal stem cells stained with DiOC18 before (A) and after (B) magnetic stimulation. The control case was subject to no magnetic stimulation. The positions of the gel disks are indicated by circles on the figure.
spacers. After 2 h, the ferrogel was cut into various shapes and placed in a large volume of distilled water, for a minimum of 24 h, to remove any residual reagents. The gels were then frozen at −20, −80, or −180 °C and lyophilized to generate macroporous ferrogels with variable pore characteristics.

**SEM Sample Preparation.** The SEM samples in Fig. 1B were prepared by simply cutting the macroporous ferrogels before hydration incubation. To prepare the SEM samples in Fig. 2D, hydrated macroporous ferrogels were frozen in the undeformed and deformed states in liquid nitrogen, and then lyophilized and sectioned for observation.

**Pore Size and Connectivity Evaluation.** The average sizes of the pores in macroporous gels were calculated by averaging the diameters of the pores in the gel. To assess the pore connectivity, the gels were soaked in a solution containing 1.0 × 10^6 μL FluorSpheres red fluorescent microspheres (Molecular Probes) with 1-μm diameter and subject to 120 cycles (on/off) of the external magnetic field. The gels were then dehydrated and examined using microcomputed tomography (micro-CT) to determine the distribution of the fluorescent beads within the gels. Micro-CT images visualized were obtained from the midplane (2.5-mm above the bottom of gels) of the cylindrical-shaped ferrogel scaffolds (5 mm in height). Fluorescent beads were imaged at high resolution (HMXST225, X-Tek; Nikon Metrology NV) with the scanner set to a voltage of 100 kV and a current of 100 A. Next, a series of 2D images of each sample was taken with an X-Tek X-ray machine while the sample was rotated 360°. The series of 2D images were converted to 3D images using CT Pro software. Image rendering and subsequent image processing including movie production were performed with Volume Graphics studio MAX. The pore connectivity was further evaluated using a water wicking technique in which the interconnected porosity was calculated as the interconnected void volume over the total volume. To determine total volume, gels were soaked in water for 1 h and weighed. A Kimwipe was then used to wick away water within interconnected pores and the gels were weighed once again. The interconnected void volume was calculated as the volume of water wiped from the gels.

**Mechanical Testing.** The nanoporous ferrogel and macroporous ferrogel were cut into cylinders (15-mm diameter, 8-mm height). The cylinders were subject to compression tests using an Instron 3342 from Instron with a strain rate of 20% per minute. Engineering stresses and strains were recorded. The ferrogel cylinders were kept hydrated throughout the tests.

**Controlled Release of Mitoxantrone.** The macroporous ferrogels in dry state were cut into disks with a volume ~2 mL. Each disk was allowed to absorb 1 mL of an aqueous solution of mitoxantrone with a concentration of 300 mg/mL. The resulting gels were stored at room temperature for ~6 h. The macroporous ferrogels were then placed in a PBS solution. Every 30 min, one group of the macroporous ferrogels was subjected to 120 cycles (on/off) of the magnetic field over 2 min, while no magnetic field was applied to the other gels. The medium surrounding each disk was collected and replaced with fresh buffer every 2 h and after each period of magnetic stimulation. To determine the concentration of plasmid DNA released into the medium was measured using QuantiTm™ PicoGreen® dsDNA Assay Kit (Invitrogen).

**Controlled Release of SDF-1α.** Macroporous ferrogels with a volume ~1 mL were allowed to absorb 0.5 mL of an aqueous solution of human SDF-1α (PeproTech) with a concentration of 2 μg/mL. Ferrogel disks were used for the release study following a procedure similar to that described in the controlled release of mitoxantrone, except the magnetic field was applied every 2 h instead of every 30 min. The medium surrounding each disk was collected and replaced with fresh buffer every 2 h and after each period of magnetic stimulation. SDF-1α protein concentrations were then determined using an Enzyme Linked Immunosorbent Assay Kit for human SDF-1α (R&D Systems).

**Controlled Cell Release in Vivo.** Mouse dengue ferrisferrol (1.5 × 10^6) (Lonza) were seeded in each ferrogel gel disk, and the gel disks were incubated at 37 °C for 4 h to allow cell adhesion. Thereafter, the ferrogel disks were placed in FBS-supplemented DMEM and the cells were released following a similar procedure as described in controlled release of mitoxantrone, except the magnetic field was applied every 2 h instead of every 30 min. The medium surrounding each disk was collected and replaced with fresh DMEM every 2 h and after each period of magnetic stimulation. The numbers of cells released were counted with a Z2™ Coulter Counter (Beckman Coulter). The cells released by magnetic stimulation were also incubated in a Petri dish with DMEM at 37 °C to monitor adhesion and proliferation.

**Controlled Cell Release in Vitro.** Mouse mesenchymal stem cells (1.5 × 10^6) (ATCC) stained with DIOC18 (Invitrogen) according to manufacturer’s protocol were seeded in each ferrogel gel disk, and the gel disks were incubated at 37 °C for 4 h to allow cell adhesion. The gel disks were implanted subcutaneously into the back region of female nude mice (NU/J, Jackson Laboratory) under anesthesia, and the incisions were closed by 5-0 Ethilon sutures (John- son & Johnson). In order to create a space for placement of gels and cell release, liquid pockets were generated around the ferrogels by subcutaneous PBS injection prior to gel placement. One hour after the implantation, the ferrogels were subject to 120 cycles (on/off) of the external magnetic field by approaching and retracting a magnet against the skin of the mouse. Fluorescence images of mice were obtained on a Xenogen IVIS Spectrum system (Caliper Life Sciences). Animal work was performed in compliance with National Institutes of Health and institutional guidelines.

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