Magnetic control of cellular processes using biofunctional nanoparticles

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Remote control of cellular functions is a key challenge in biomedical research. Only a few tools are currently capable of manipulating cellular events at distance, at spatial and temporal scales matching their naturally active range. A promising approach, often referred to as ‘magnetogenetics’, is based on the use of magnetic fields, in conjunction with targeted biofunctional magnetic nanoparticles. By triggering molecular stimuli via mechanical, thermal or biochemical perturbations, magnetic actuation constitutes a highly versatile tool with numerous applications in fundamental research as well as exciting prospects in nano- and regenerative medicine. Here, we highlight recent studies, comment on the advancement of magnetic manipulation, and discuss remaining challenges.

1 Introduction

Remote activation of cellular processes constitutes an important challenge in nanotechnology and bioengineering. This challenge can now be addressed with techniques based on advanced physical (optical, magnetic or electrical) and chemical modalities which go well beyond conventional genetic or pharmacological approaches. These novel techniques enable systematic activation at the single cell level but within environments as complex as a living organism. Recent studies, for example, succeeded in the control of oriented migration, intracellular transport, gene expression, and differentiation. Thus, it becomes possible to probe specific biological mechanisms, to quantitatively interrogate the complex molecular circuitries that mediate signaling events, or even to control cell behavior for regenerative medicine applications.

Optical techniques based on photocontrol of protein activity, using photoactivatable reagents or optogenetics, have been very beneficial, with a broad range of applications in cell biology and neuroscience. Yet, light-based methods also present some limitations, such as the need to express multiple genetically-modified photo-reactive proteins, the potential phototoxicity of optical stimuli in the biological specimen, or the difficulty to apply sustained spatially-patterned illumination, which can be challenging in thick and scattering tissues.

An emerging and complementary approach to remotely control biological processes is based on magnetic stimulation. So far, the effect of magnetic fields on cellular response has been investigated in different contexts. A first instance is the field of ‘magnetosensation’, i.e. the ability of some organisms to detect magnetic fields, e.g. for the purpose of navigation. A second area of research focuses on the influence of magnetic fields on biological processes in general. Here, usually strong magnetic fields (>1 T) have been applied to show that biopolymers with high diamagnetic anisotropy (e.g. microtubules and nucleic acid chains) can respond to the external magnetic field. While both of these fields are active and interesting areas of research, this review elaborates on a third approach, in which the magnetic control of biological processes is mediated by functionalized magnetic nanoparticles (MNPs). Recent studies have indeed demonstrated how MNPs can serve to convert the external signal of a static or oscillating magnetic field of moderate strength (~100 mT, field gradient ~10¹⁸ to 10¹⁹ T m⁻¹) into biological events (see Table 1 for an overview). These studies, which go well beyond the conventional use of MNPs as imaging contrast agents or drug-delivery carriers, demonstrated how different magnetic field conditions effectuate mechanical, thermal or biochemical stimuli that trigger a specific cellular event. The benefits of such flexibility are further reinforced by the fact that magnetic fields can non-invasively penetrate deep into tissues, wherefore cellular functions and circuits can be probed in live cultured cells, tissues and organisms. Altogether, these novel manipulation tools have laid the ground for a new field, sometimes termed ‘magnetogenetics’. Here, we review recent advances in magnetogenetics and highlight its potential for both fundamental and applied biomedical research.

2 Magnetic control of cellular processes

Magneto-mechanical stimulation

The use of MNPs in cell biology has long been associated with the study of mechano-transduction, that is, the conversion of...
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**Table 1** Overview of recent magnetic manipulation studies and reference values

<table>
<thead>
<tr>
<th>Application</th>
<th>Mechanism</th>
<th>Force $F$/magnetic flux density $B$/amplitude magnetic flux density $B_0$/field gradient $\nabla B$</th>
<th>MNP</th>
<th>Reference</th>
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<tr>
<td><strong>Magneto-mechanical stimulation</strong></td>
<td>Stretching/pulling by magnetic tip</td>
<td>$F = 1$–$47$ pN</td>
<td>Core: 10–30 nm zinc-doped iron oxide, coating: silica &amp; gold shell; thiolated DNA</td>
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<td>Control of Notch &amp; E-cadherin receptor activity</td>
<td>Stretching/twisting by 3D magnetic multipoles</td>
<td>$B \sim 250$ mT, $B &lt; 2.5$ mT (twisting field)</td>
<td>Core: 4 μm ferromagnetic bead, coating: RGD peptides</td>
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<td>Stretching of chromatin for gene transcription upregulation</td>
<td>Pulling/clustering by electromagnetic (EM) or permanent magnet (PM)</td>
<td>$B = 50$ mT (EM &amp; PM), 500 mT (PM)</td>
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<td>Control of TRPV1 ion channel gating</td>
<td>Stretch/pulling by oscillatory motion on magnetic arrays</td>
<td>$B = 25$–$120$ mT</td>
<td>Core: 300 nm iron oxide, coating: antibodies or RGD tripeptide</td>
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<td>Control of Wnt-frizzled receptor activity</td>
<td>Attraction/pulling by magnetic array</td>
<td>$F = 100$ nN, $B = 25$–$100$ mT, $\nabla B = 2500$–$7000$ T m$^{-1}$</td>
<td>Coating: dextran</td>
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<td>Stimulation of filopodia formation &amp; oriented cell division</td>
<td>Attraction/pulling by magnetic tip</td>
<td>$F = 0.1$ pN, $\nabla B = 10^3$ T m$^{-1}$</td>
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<td>Magnetic tweezer induced mechanical tissue deformation</td>
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<td>Core: 7.5 nm magnetite, coating: citrate molecules</td>
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<td>Control of Drosophila embryonic tissue deformation for gene expression</td>
<td>Pulling/clustering by permanent magnet</td>
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<td>Modulation of cell endocytosis</td>
<td>Heat activation via radio-frequency waves</td>
<td>$f = 40$ MHz, $B_0 = 1.3$ mT</td>
<td>Core: 6 nm manganese ferrite, coating: streptavidin &amp; PEG-phospholipids, SLP (specific loss power) = 2.5 W g$^{-1}$</td>
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<tr>
<td>Control of TRPV1 ion channel gating</td>
<td>Heat activation via radio-frequency waves</td>
<td>$f = 500$ kHz, $B_0 = 18$ mT</td>
<td>Core: 22 nm iron oxide, coating: PEG, SLP = 660 W g$^{-1}$</td>
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<td>Control of TRPV1 ion channel gating for neuronal cell excitation</td>
<td>Heat activation via radio-frequency waves</td>
<td>$f = 465$ kHz; $B_{ag} \sim 5$ mT (Stanley 2012) &amp; ~30 mT (Stanley 2015/2016); $B \sim 0.1$–$1$ T (Stanley 2016)</td>
<td>Core: 20–25 nm iron oxide (Stanley 2012), coating: carboxylic acids (Stanley 2012), endoferritin particle (Stanley 2015/2016), SAR (specific absorption rate) = 0.63 W g$^{-1}$ (Stanley 2012)</td>
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<td>Clustering by electromagnetic tip</td>
<td>$F \sim 0.01$ fN, $B \sim 100$ mT</td>
<td>Core: 5 nm iron, coating: 10 nm polymer shell, with amine groups</td>
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<td>Control of FcR1 receptor activity for inflammatory responses</td>
<td>Attraction/clustering by magnetic arrays</td>
<td>$F \sim 30$ fN, $B \sim 200$ mT (magnetic arrays), $B \sim 500$ mT (zebrafish)</td>
<td>Core: 15 nm zinc-doped iron oxide, coating: thiols</td>
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<td>Attraction/pulling by magnetic tip</td>
<td>$F \sim 10$ fN, $B \sim 150$ mT, $\nabla B = 50$ T m$^{-1}$</td>
<td>Core: 100 nm iron oxide, coating: 10 nm polymer shell with carboxylic acids</td>
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<td>Attraction/pulling by magnetic tip</td>
<td>$F \sim 10$–$30$ pN, $B \sim 200$ mT, $\nabla B = 10^3$ to $10^4$ T m$^{-1}$</td>
<td>Core: 500 nm iron oxide, coating: 10 nm polymer shell with streptavidin</td>
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<td>Control of intracellular Rac-GTPase signalling</td>
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a mechanical stimulus into an electrical or biochemical signal.25

Attaching a bead at a cell surface to bend, stretch, or twist the cell membrane has proven very useful to analyze cell mechanical features26–28 and the transduction of mechanical constraints by membrane-associated molecular complexes, such as integrins29–32 or cadherins.27,30 In recent years, the scope of these experiments has been largely expanded by means of carefully designed and targeted MNPs.33–35 Combined with advanced magnetic systems, these MNPs enabled the manipulation of various biological processes with spatio-temporal resolution in the submicrometer and millisecond range, and down to the level of single proteins.34

One such example is the work by Seo et al.,7 where monofunctionalized iron-oxide nanoparticles capped with a gold shell (with ~50 nm total diameter) were used to manipulate individual Notch receptors or E-cadherin at the cell surface. Utilizing a calibrated tweezer setup, they found that individual Notch receptors require a critical loading force in the 2–9 pN range to activate downstream signaling. For E-cadherin, they identified the importance of cooperative spatial aggregation and mechanical loading to achieve actin and vinculin adaptor recruitment (Fig. 1a).

Remote actuation of signaling by magneto-mechanical stimulation may also be realized through direct gating of ion channels via deflection or stretching. For instance, Lee et al.15 demonstrated ultrafast (sub-ms) mechanical control of the deflection-relaxation dynamics of stereocilia bundles bound to cubic MNPs (size ~50 nm) and submitted to oscillating fields in inner ear hair cells (Fig. 1b). Wheeler et al. used the ‘transient receptor potential channel subfamily V member 4’ (TRPV4), a membrane receptor known to gate in response to osmotic, chemical or mechanical cues.36–39 By fusing TRPV4 to genetically-encoded ferritin nanoparticles and applying a static magnetic gradient, they triggered Ca2+ transients, which elicited action potential in neurons, enhanced tactile behavior in zebrafish, and affected reward behavior in mice.39 It should be noted though that the physical mechanism underlying the actuation of TRPV4 by endogenous ferritin is a subject of current debate (see discussion).

Other studies have aimed at controlling developmental processes, such as stem cell differentiation or tissue organization.7,40–43 For instance, transmembrane ion channels TREK-1, which are responsible for setting the resting membrane potential and intracellular Ca2+-concentrations, and which influence differentiation processes, were labeled in human mesenchymal stem cells with 250 nm MNPs and were stimulated by applying a slowly oscillating (~1 Hz) magnetic field over a period of 7 or 21 days in vitro.7 The resulting increase in the proteins Sox9, osteopontin and collagen provided evidence that a mechanical stimulation of TREK-1 is sufficient to induce the differentiation of osteoprogenitor cell populations toward an osteogenic lineage. Next to the overall change in gene expression, an interesting point raised by these studies is that long-term magnetic stimulation over a day and up to several weeks showed a significant effect on the targeted gene expression. The expression of the reporter molecules of TREK-1, for example, exhibited a 2-fold increase after 7 days of magnetic field exposure7 and a similar increase was observed in a study of the Wnt signaling pathway, where expression levels were compared after 6 h and 1 day.46 Hence, investigating the temporal effects of magnetic stimulation is important to achieve full control of a biological process.

Desprat et al. demonstrated how mechanical forces during Drosophila embryo development are coupled to the regulation of TWIST, a transcription factor vitally involved in early anterior endoderm cell differentiation;44 first, the natural compression movement occurring at the onset of Drosophila gastrulation was suppressed and TWIST expression remained low. Thereafter, the natural deformation of cells was mimicked by pulling on ferrofluid loaded cells in the neighborhood of the otherwise naturally compressed cells. This mechanical stimulus rescued TWIST up-regulation and resulted in gene expression profiles similar to those occurring naturally (Fig. 1c). More recently, a similar approach was used to show how mechanical pressure contributes to the onset of tumorigenesis in a mouse model for colon tumor development.45

Finally, directional magnetic pulling of MNPs internalized in endosomes was used by the group of Di Carlo in different biological contexts – e.g. to induce coordinated filopodia formation, to bias spindle orientation in dividing cells,46,47 as
well as to mechanically stimulate calcium influx and polarity of neurons. A breakthrough in these studies was to establish a massively parallelized assay, where microfabricated magnetic arrays, with cells plated in close proximity to one micromagnet, enabled high-throughput measurements with unprecedented control. With this assay, the group provided an elegant solution for one of the major challenges in many magnetic manipulation studies, namely, the application of well-defined magnetic forces over a large population of cells.

**Magneto-thermal stimulation**

A second magnetic actuation modality is based on the thermal response of magnetic nanoparticles. When placed in a radio-frequency (~1 MHz) magnetic field, some MNPs (depending on their size, shape, and magnetic content) are able to convert the field stimulation into heat. Thus, they can be used as local hotspots to stimulate thermo-responsive molecules – several of which are naturally found in mammalian cells. The most prominent examples are proteins of the ‘transient receptor potential’ (TRP) channel family, ion channels which are located in the plasma membrane of cells. TRPM8 and TRPA1, for example, are ion channels sensitive to cold temperatures (active between 25–28°C and <17°C, respectively) and TRPV1, TRPV3, and TRPV4 (active >42°C, >38°C and >35°C, respectively) are activated when heated. Note, that the most suitable activation temperature can vary between different cell types due to their different thermosensitivity. The studies discussed in the following, report the control of TRPV1 cation gating in different cells and this gating in turn was used to trigger specific signaling pathways (Fig. 2a).

The magneto-thermal stimuli reported in recent studies enabled the control of molecular activity states in single cells, tissue and animals. More precisely, calcium influx into cells was controlled via TRPV1 gating and evidenced at the single cell level, either in HEK293 cells or in hippocampal neuronal cells leading to the firing of action potentials. In animals, magneto-thermal actuation was first applied in *C. elegans* worms where heating of MNPs near sensory neurons triggered a thermal avoidance, i.e. a spatial retraction response. In mice, activation of TRPV1 in ventral tegmental brain regions evoked considerable neuronal excitation (Fig. 2b) and highlighted the potential of magneto-thermal tools for remote and local deep-brain stimulation without the need for implants and connectors. The remote gating of TRPV1 was achieved by means of a high concentration (~mg ml⁻¹) of synthetic iron-oxide MNPs (size 10–30 nm) either freely diffusing in the cell cytoplasm or targeted to the cell plasma membrane via biotin–streptavidin interactions to increase the efficiency of ion channel heating as well as to reduce side effects.

In a different set of experiments, Stanley *et al.* followed a similar strategy in order to control insulin expression and, consequently, the level of blood glucose in mice (Fig. 2c). To this end, they used endogenous ferritin as heat generating nanoparticles targeted to TRPV1 to thermally gate the ion channel, and thus trigger TRPV1-mediated calcium influx.

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**Fig. 1** Magneto-mechanical stimulation. (a) Left: Mechanoreceptor activation via pulling. A magnetic tip in close proximity of an MNP bound receptor mechanically loads the MNP–receptor complex and activates intracellular signaling. Right: Immunofluorescence staining for MNPs, E-cadherin and the signal activation reporters, actin and vinculin. 9 pN forces result in enhanced molecular colocalization and recruitment compared to 1 pN forces. Scale bar, 2 μm. Taken from Seo *et al.* with permission from Elsevier. (b) Left: Gating ion channels of the inner ear via mechanical deflection. A magnetic field gradient tilts stereocilia with attached MNPs. Bundle displacement opens the tip link at the channel top (yellow) resulting in ion influx. Right: Traces of pulsed magnetic stimulation, the corresponding hair bundle displacement, the fluorescence signal of a Ca²⁺-sensitive dye inside the bundle, and the control experiment, where channel opening was prevented. Reprinted with permission from Lee *et al.* Copyright 2014 American Chemical Society. (c) Left: TWIST expression correlates with mechanical cell compression in a *Drosophila* embryo. To probe this, the natural compression of endodermal cells is blocked and mimicked instead by magnetically pulling on cells containing a ferrofluid. These magnetized cells are adjacent to endodermal cells and compress the tissue upon magnetic field application. Right: (i) cell compression induced by magnetic manipulation (yellow arrow). (ii) Particle imaging velocimetry indicates compression changes – lowest in black and highest in red. (iii) Inhibition of TWIST expression in the uncompressed cells of the ablated embryo (between the red arrows). (iv) Recovery of TWIST expression in the cells of the ablated embryo by inducing physiological compression with magnetic fields. Taken from Desprat *et al.* with permission from Elsevier.
Fig. 2  Magneto-thermal stimulation. (a) Controlling membrane ion channels via MNP heating. An MNP in close proximity to the temperature-sensitive ion channel TRPV1 is locally heated to >42 °C with a radio-frequency (RF) magnetic field. Subsequently, TRPV1 opens, enabling Ca2+ influx. The ion concentration increase is used to activate selected signaling processes. (b) Magneto-thermal TRPV1 stimulation for the controlled activation of hippocampal neurons. The ion channel gating evokes correlated and repeated trains of action potentials in TRPV1 expressing neurons. Top: 10 fluorescence traces (orange) with an average overlay (black) before, during, and after magnetic stimulation (blue bar). Bottom: Raster plots of 100 randomly selected neurons exhibiting repetitive calcium spikes. Shaded blue bars represent alternating magnetic field pulses. Taken from Chen et al.56 with permission from AAAS. (c) Remote regulation of glucose homeostasis in mice by TRPV1 activation. During channel activation in a RF magnetic field (pink area) a calcium-dependent transgene expression of insulin is initiated. Enhanced insulin expression followed by reduced blood glucose levels in mice are predominantly observed for ferritin–MNPs directly coupled to TRPV1 via the GFP–antiGFP nanobody interaction (sGFP–TRPV1/GFP–ferritin), as well as for ferritin–MNPs associated to the cell plasma membrane (TRPV1/myrferritin). *P < 0.05. Data are mean ± s.e.m. Taken from Stanley et al.57 with permission from AAAS.

Magneto-molecular stimulation

A third magnetic actuation modality relies on the control of biomolecular activity patterns in the cell by modulation of the concentration and spatial distribution of signaling molecules. An early example is the use of MNPs specifically bound to membrane receptors, to control the oligomerization-dependent activation mechanisms in the cell plasma membrane.59-61 Here, a static field is applied to reversibly cluster receptor–MNP complexes through dipole–dipole interactions between the nanoparticles, which subsequently triggers an intracellular signaling response. By doing so, Mannix et al. probed the immune surveillance as performed by mast cells and found that aggregation of FccRI–dinitrophenyl receptor–ligand complexes was sufficient to increase cytosolic calcium and to initiate a local inflammatory response69 (Fig. 3a). Similarly, the magnetically-induced dimerization of EGF receptors led to the activation of downstream signaling cascades, as evidenced by the phosphorylation of the receptors.68 Building on this approach, Cho et al. targeted iron-oxide MNPs to the death receptor 4 (DR4) of colon cancer cells to mimic the natural apoptosis signaling pathway via MNP-DR4 aggregation in vitro. When injected into a zebrafish embryo, these MNPs targeted a DR4 related receptor and aggregated upon magnetic field stimulation in the zebrafish tail, altering its morphology.64

Besides the ability to activate signaling pathways at the plasma membrane receptor level, magnetogenetics also permits the investigation and control of the intracellular signaling machinery, or, more generally, of the subcellular organization. This approach faces the challenges of nanoparticle internalization in the cytosol (escaping endosomal pathways) and surface passivation to ensure colloidal stability inside the complex cytoplasmic environment. In the case of subcellular manipulation, MNPs tagged with signaling proteins and in the fluid phase of the cytoplasm can function as signaling nanostructures, able to interact with downstream effectors and to activate signaling pathways. With magnetic gradients, these nanostructures can further be displaced through the cytoplasm and accumulate at specific locations in order to modulate the local concentration of signaling molecules.62,63 Etoe et al. used this approach with 500 nm-MNPs coupled to Tiam1, a GEF molecule for the small GTPase Rac1.64 Once brought to the plasma membrane, MNPs triggered actin cytoskeleton remodeling and cell protrusion formation (Fig. 3b). Another experiment in droplets of Xenopus egg extract probed the spatial arrangement of the Ran/RCC1 signaling pathway involved in cell cytoskeleton and mitosis regulation.65 When accumulating the particles in a field gradient, RanGTP coupled MNPs were found to act as a bio-inspired switch where microtubule aster growth is only
initiated if a distinct concentration threshold is exceeded (Fig. 3c).

3 Conclusion and perspectives

All of the above examples attest to the great variety of biological questions that can be addressed using magnetic manipulation mediated by MNPs (see Table 1 for an overview). A key factor is the unique spectrum of magnetic actuation modalities (mechanical, thermal, and biochemical) by which quantitative and spatiotemporally modulated stimuli can be generated.

So what can be expected from magnetogenetics in the future? An exciting prospect is the possibility to control cellular behavior and tissue engineering directly in organisms where, contrary to light illumination, magnetic fields can penetrate easily and without inducing damage. Conceivably, remote activation of cellular processes, such as cell differentiation or oriented migration, might contribute to the success of cell-based therapies, for instance to target and differentiate stem cells at sites of injury, during lesion repair, or for the treatment of neurological disorders. Magnetic stimulation might also become a valuable and non-invasive technique for the control of hormone release and metabolic activity during pathologies.

Interestingly, MNPs can serve to hijack signalling pathways and exploit the ability of individual cells to sense signalling cues with great sensitivity, to process signalling information (including its amplification), and to mobilize complex molecular machineries to ensure a proper response. Thus, magnetic stimuli provide a means to non-invasively instruct cells to carry out tasks (such as differentiation and growth) that they are already programmed to perform, but do not necessarily accomplish at the right time and the right place. In view of realizing such control of vital biological functions, magnetogenetics will also benefit from further studies in the fields of ‘magnetosensation’ and magneto-responsive biomolecules, where the effect of magnetic fields in the absence of MNPs is investigated.

While much work is still needed before magnetogenetics becomes a versatile tool in the arsenal of regenerative medicine and nanomedicine, several stimulating applications can already be envisaged in the short term. First, the ability to displace or heat MNPs at sub-micrometer scales, and thereby to apply forces, activate molecular functions or trigger biochemical reactions, should prove invaluable to get novel insights into the mechanisms of biological processes. Second, as is well known in engineering, applying a variable input and measuring the output is a powerful means to dissect the molecular circuits underlying the cellular response to external cues. By quantitatively modulating the amplitude, frequency, and spatial localization of the stimulus, one can identify key aspects of the cue processing, such as amplification, filtering or the existence of thresholds. These important systems-level features are usually difficult to infer from the genetic manipulation of molecular components. Finally, magnetic manipulation involving MNPs might serve for the design of (semi-)synthetic molecular machines inside the cell, i.e. the engineering of biological machines inside the cell, i.e. the engineering of biological
oscarators and switches based on MNPs coupled to molecular
nanoplatforms.

Several substantial challenges remain to take full advantage of
the promises of magnetogenetics. Novel MNPs will surely be
needed, with physical [size and shape], chemical [composition] and
biofunctional surface properties tailored to ensure optim-
ized magnetic response and efficient biotargeting as well as
passivation against non-specific interactions. An obvious chal-
lenge and well-identified issue in bionanosciences is the deliv-
ery of MNPs within cells.66 Here, development of cell
loading protocols alternative to injection, endocytosis, or per-
meabilizing reagents is needed, in particular, when anticipat-
ing magnetic manipulation with MNPs inside organisms.

A solution might reside in the use of genetically-encoded
MNPs, in the form, for instance, of natural ferritin cage
proteins67–71 or viral capsids.36,57,72 Yet, a difficulty with
endogenous ferritin is that they store iron as ferrihydrite, which
has reduced magnetic properties compared to iron oxides.77–78
Several strategies can be envisaged to enhance the magnetic
response of fully genetically encoded systems. For ferritin-
expressing eukaryotic cells, Kim et al.73 established protocols
to elevate intracellular iron levels by combining ferritin
expression with DMT1-based iron import and ferrous ammo-
nium sulfate-supplemented culture medium. Another approach
might be to exploit the ability of magnetotactic bacteria to
synthesize biomineralized Fe3O4 crystals, so-called magne-
tosomes. Recently, Kolinko et al. have transferred this bio-
mineralization potential from magnetotactic bacteria to a
foreign non-magnetic prokaryotic organism.73 This successful
deed is an exciting step towards the more challenging goal
of enabling eukaryotic cells to synthesize tailored magnetic
nanostructures.

To increase the magnetic response, a complementary
strategy is to improve our ability to generate strong magnetic
gradients.74 Yet, the main difficulty is that applying strong
gradients implies working at a short distance, which is partic-
ularly challenging in tissues and organisms where distances
between the magnetic device and the nanoparticles are in the
millimeter/centimeter range. One interesting way to overcome
this challenge might be the use of locally magnetizable
implants.75–76

Importantly, the development of magnetogenetics requires
a careful and quantitative examination of the physical mecha-
nisms of action. In particular, the activation mechanisms
proposed in some of the recent studies, notably using endoge-
 nous ferritin nanoparticles,69–70 have been heavily questioned.
Indeed, based on physical arguments and the experimental
conditions reported, it was estimated that the force and/or
torque exerted by the endogenous ferritin NP on the attached
TRPV4 channel are 4 to 9 orders of magnitude lower than those
due to thermal agitation.88 Hence, it is difficult to comprehend
how mechanical actuation could be achieved. Similarly, the
heating response in a RF field, even with a ferritin optimized
with cobalt doping, is about ten orders of magnitude too
small.88 Thus there is a need for clarifying the exact nature of
the physical or biochemical stimuli induced by the magnetic field
and MNPs in these experiments.69–70

To conclude, we anticipate that, in view of its benefits as well
as numerous challenges and open questions, the magnetic
control of cellular behavior will become an active field of
research in the forthcoming years. With its successful advan-
cement, magnetogenetics should be established as a key
technology, complementary to optogenetic tools, and will find
multidisciplinary applications in chemistry, bioengineering,
bionanosciences, biology, biophysics, or neurosciences.

Conflicts of interest
There are no conflicts of interest to declare.

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