

1 Automatic navigation of an untethered device in the artery of a living 2 animal using a conventional clinical magnetic resonance imaging system

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22 The feasibility for *in vivo* navigation of untethered devices or robots is demonstrated with the
23 control and tracking of a 1.5 mm diameter ferromagnetic bead in the carotid artery of a living swine
24 using a clinical magnetic resonance imaging (MRI) platform. Navigation is achieved by inducing
25 displacement forces from the three orthogonal slice selection and signal encoding gradient coils of
26 a standard MRI system. The proposed method performs automatic tracking, propulsion, and
27 computer control sequences at a sufficient rate to allow navigation along preplanned paths in the
28 blood circulatory system. This technique expands the range of applications in MRI-based
29 interventions. © 2007 American Institute of Physics. [DOI: 10.1063/1.2713229]
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31 The replacement of many highly invasive procedures by
32 minimally invasive surgeries (MIS) has been a significant
33 milestone for modern interventional medicine.¹ Although the
34 next step in modern interventional procedures² may be in-
35 spired from the 1966 science fiction movie “Fantastic Voy-
36 age,” forty years later, techniques allowing automatic navi-
37 gation and directed targeting of untethered objects, devices,
38 or microscale robots in deep regions of the human body have
39 not been realized yet.³ Medical interventional procedures re-
40 lying on untethered instruments, carriers, or sensors navigat-
41 ing in the bloodstream could open the path to a variety of
42 therapies, interventions, and diagnostic methods directly tar-
43 geted at operative sites that remain inaccessible or at high
44 risk to modern MIS-based instruments. In larger blood ves-
45 sels, navigation of larger untethered devices may assist in
46 applications such as reopening of encumbered arterial ways
47 or alternative methods of aneurysm treatment, to name but a
48 few applications. Devices released from a catheter could per-
49 form as an extension of existing MIS-based instruments.

50 Here, a conventional magnetic resonance imaging (MRI)
51 system as found in many clinical settings is considered. Al-
52 though MRI systems have been used exclusively for imaging
53 purposes until now, we show that its range of potential medi-
54 cal and research applications can be expanded beyond imag-

ing by this demonstrated ability of MRI scanners to provide 55
a means of propulsion for minimally invasive devices. This 56
actuation method combined with imaging/tracking and con- 57
trol software represents the components essential for precise 58
navigation of various types of untethered interventional de- 59
vices and objects inside the human body. The use of a clini- 60
cal MRI system for such applications offers many advan- 61
tages including but not limited to enhanced tissue contrast, 62
lack of radiation, and widespread availability in clinical en- 63
vironments. It is also shown that for navigation in larger 64
blood vessels, the three orthogonal gradient coils inside the 65
bore of an MRI system, typically used for slice selection and 66
signal encoding during MR imaging, can also induce a three- 67
dimensional (3D) directional magnetic force sufficient to 68
propel an object made of ferromagnetic material.⁴ This force 69
is proportional to the amplitude of the gradient vector ac- 70
cording to 71

$$\mathbf{F} = RV(\mathbf{M} \cdot \nabla)\mathbf{B}, \quad (1) \quad 72$$

where R is the duty cycle of gradient generation, V the vol- 73
ume of the ferromagnetic object, \mathbf{M} the magnetization of the 74
material, \mathbf{B} the magnetic field, and ∇ the vector differential 75
operator. To navigate a spherical device in the center of 76
larger blood vessels with Reynolds number $1000 < \text{Re}$ 77
 < 3000 and considering the retarding effect of the blood ves- 78
sel walls,^{5,6} magnetic propulsion force must overcome a drag 79
force⁷ 80

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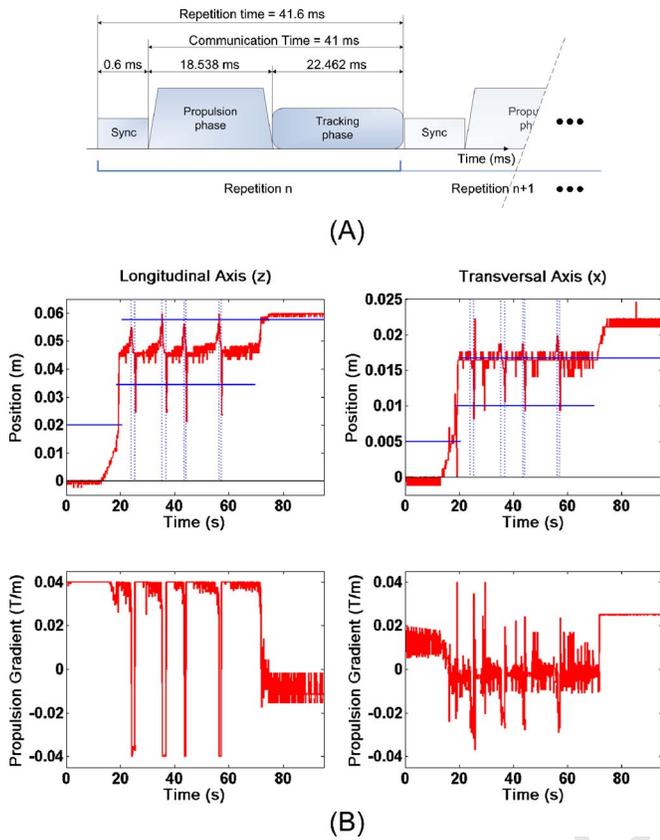


FIG. 1. (A) Overview of the real-time navigation MRI sequence for the displacement and control of the magnetic sphere. (B) *In vivo* control of a 1.5 mm ferromagnetic sphere along 11 waypoints inside the carotid artery of a living swine: tracking and propulsion gradient histories. In (a) and (c) time varying line shows sphere position along the corresponding axis. Horizontal lines represent waypoints 1, 2, and 3 targeted by the magnetic sphere. Vertical dotted lines indicate a target waypoint change during oscillation phase. (b) and (d) show gradient amplitudes as a function of time. The Y axis is not displayed for clarity since no vertical control gradients was required during this particular experiment.

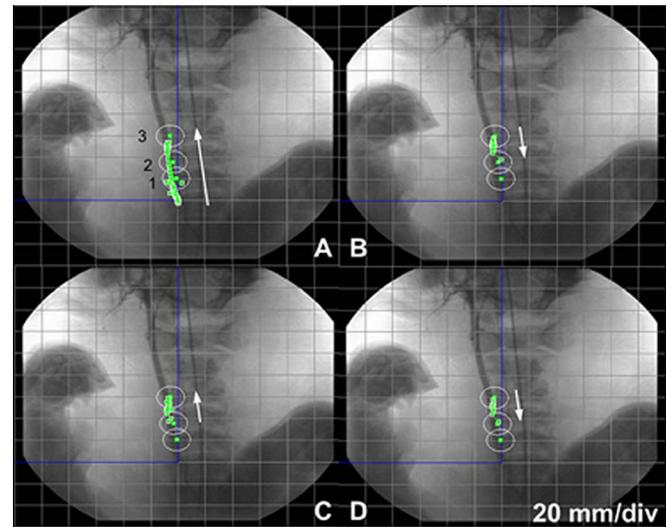


FIG. 2. *In vivo* automatic navigation of a 1.5 mm ferromagnetic bead inside the carotid artery of a living swine. The trajectories are superimposed over an x-ray angiography. The line of dots over the artery shows actual displacement of the bead. The circles (20 mm diameter) around each waypoint show precision tolerance region. Arrows show the direction of displacement. Image A, the bead travels through waypoints 1 and 2 and reaches the precision region of waypoint 3. Target waypoint changes from waypoint 2 to waypoint 3 (image B). Images C and D show two subsequent round-trips as target waypoint switches from waypoint 2 to waypoint 3 alternatively.

running, a processing time and a communication time are required for the control routine to be completed and the command to be sent to the running sequence, setting the minimum duration of the latter event with a maximum duration set by power limitations from the gradients applied during the tracking and propulsion phases. A longer duration provides more magnetic propulsion force but increases the loading level on the gradient coils and amplifiers. Hence, the 41 ms duration was determined to be the maximum level of power dissipation the MRI scanner used for the experiment could sustain.

A ferromagnetic 1.5 mm (0.0136 g) diameter chrome steel sphere (Salem Specialty Balls Company, Canton, CT) was used for the experiment. Its magnetization $M_{1.5 T} = 1.35 \times 10^6$ A/m was measured at $B_0 = 1.5$ T (Walker Scientific VSM, Worcester, MA) being the field present inside the bore of the MRI system used (Siemens Magnetom Avanto 1.5 T, Erlangen, Germany). A spherical shape was chosen instead of a prolate spheroidal shape that minimizes drag force⁸⁻¹⁰ since a ferromagnetic body tends to maintain its original orientation along the B_0 field inside the MRI bore. Hence, the symmetry of a spherical body although not optimal for a predetermined direction of motion is still a suitable and simple compromise for general two-dimensional and 3D navigations.

The tests were performed with real-time MRI positional accuracy of 0.59 mm combined in a time multiplexed fashion with propulsion using the MRI imaging gradient coils and proportional-integral-derivative control. Real-time tracking bypassing MR susceptibility artifacts¹¹ caused by the ferromagnetic material was performed.

A tracking data history during *in vivo* navigation of the ferromagnetic sphere being automatically controlled inside the carotid artery of a 25 kg living swine (Supplementary Movie 1, see Ref. 13) is depicted in Fig. 1(b) and Fig. 2. Ten round-trips were programmed and achieved during this test.

$$|\mathbf{D}| = \frac{1}{2} \rho \left(\frac{u_t}{1 - \lambda^{1.5}} \right)^2 A C_{D\text{sphere}}, \quad \lambda < 0.6, \quad (2)$$

where ρ is the density of blood, A the frontal area of the immersed body, u_t the terminal velocity of the device, λ is the ratio between the spherical device's diameter d and the vessel's diameter, and $C_{D\text{sphere}}$ the drag coefficient,

$$C_{D\text{sphere}} \approx \frac{24}{\text{Re}} + \frac{6}{1 + \sqrt{\text{Re}}} + 0.4, \quad 0 \leq \text{Re} = \frac{\rho u d}{\mu} \leq 2 \times 10^5, \quad (3)$$

where u and μ are the relative velocity between the immersed device and the blood and the viscosity of blood, respectively.

Endovascular navigation made possible by integrating propulsion and tracking events within control software that deals with their time sequencing as well as with real-time, physiological, and technological constraints is presented in Fig. 1(a). The navigation sequence coordinated through custom developed proprietary software modules embedded in the MRI system consists of two events: a synchronization event responsible for the controller routine to be called that lasts for 0.6 ms and a propulsion phase and tracking acquisition that lasts for 41 ms. While the latter event is

137 In Fig. 1(b), the tracking history of the sphere as well as the
138 amplitude of the propulsion gradients in the coordinate sys-
139 tem of the MRI platform are shown as a function of time.

140 In addition to the trade-off between the refresh rate and
141 the duty cycle of the propulsion gradients [Fig. 1(a)], the
142 time duration (repetition time) of the tracking sequence is
143 another constraint that needs to be taken into account for
144 optimal navigation performance. For more precise navigation
145 subjected to perturbations, higher refresh or control rates are
146 preferred. On the other hand, control rates beyond some
147 thresholds can decrease tracking and propulsion performance
148 significantly. Because of MRI signal considerations, hydro-
149 gen protons require some time duration between two subse-
150 quent rf excitations to reach an equilibrium condition. With
151 insufficient repetition time, MRI signal acquisitions can be-
152 come unusable for tracking. As a general rule, repetition time
153 can be decreased until MR signal quality is degraded but still
154 usable for tracking within acceptable navigational param-
155 eters such as control rate and propulsion time especially
156 when subjected to a pulsatile flow and body movement per-
157 turbations. During the experiments, a positive 13° inclination
158 of the carotid arteries with respect to the horizontal X-Z
159 plane of the MRI system was measured from a 3D MR an-
160 giography. When coupled with a potential very low friction
161 coefficient¹⁴ between the bead and the vessel, displacements
162 of the bead when not subjected to propulsion force could
163 occur. To reduce potential undesirable motions, a higher re-
164 fresh rate of 24 Hz (41.6 ms repetition time) was chosen in
165 this particular context to guarantee stable navigation while
166 preserving slightly noisy but usable MRI tracking signals.

167 In our experimental realization of a navigable device in
168 larger blood vessels, it is shown that automatic trajectory
169 control of a ferromagnetic body is feasible without modify-
170 ing the hardware of clinical MRI systems. Here, peak veloci-
171 ties in the range of 8.43–11.1 cm/s inside the artery (5 mm
172 lumen) without blood flow have been recorded. The experi-
173 mental result is less than the theoretical value of ~ 16 cm/s
174 estimated from the previous equations using $R=44.5\%$ [Fig.
175 1(a)]. This difference may be caused by the presence of the
176 balloon catheter (used to stop blood flow in the artery) sheath
177 as well as by the eccentricity of the sphere in the vessel that
178 is not taken into account in the wall effect correlation.⁵

179 Increasing $\lambda=0.3$ to an optimal value (close to $\lambda=0.42$)
180 would improve performance by providing the best trade-off
181 between propulsion force and retarding effect caused by the
182 vessel walls.⁵ This indicates that the design of subsequent
183 devices would not only be guided by the types of interven-
184 tional procedures but by the diameter and blood flow of the
185 vessels used to reach the target. In turn, complex pathways
186 involving various types of blood vessels will influence fur-
187 ther the design of the device that could be made of a single
188 or an agglomeration of smaller biodegradable polymeric fer-
189 romagnetic entities.

190 For smaller vessels, since magnetic force scales down at
191 a cubic rate while drag force decreases linearly at low Rey-
192 nolds numbers, the addition of propulsion dedicated gradient
193 coils in the MRI bore for inducing propulsion force on a
194 smaller device must be considered. Our preliminary data ac-

quired in this demonstration suggest that due to limits in **195**
 gradient amplitudes that can be generated, the same principle **196**
 but for the purpose of steering instead of propelling devices **197**
 in low Reynolds hydrodynamic environments (capillaries to **198**
 small arterioles or venules) could be applied. On the other **199**
 hand, nanocarriers due to their single domain magnetic prop- **200**
 erty behave as MRI contrast agents that already proved to be **201**
 visible and traceable with MRI without causing image arte- **202**
 facts as it is the case for larger magnetic entities. Gradient **203**
 induced magnetic forces from an MRI platform could steer **204**
 agglomerations of micro- or nanodevices (or more complex **205**
 devices such as nanorobots in a farther future) being drifted **206**
 by blood flow. Improved targeting could be enhanced further **207**
 by decreasing blood velocity with an appropriate method **208**
 such as upstream balloon catheter inflation. **209**

The results and the methods developed for this letter **210**
 provide valuable data for potential future development of **211**
 enhanced MIS-based interventional systems and procedures **212**
 that can be adapted to a variety of applications particularly **213**
 for direct delivery in nanomedicine where extensive research **214**
 efforts are being made to develop multifunctional magnetic **215**
 nanoparticles.¹⁵ **216**

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 ed via a direct link in the online article's HTML reference section or via **249**
 the EPAPS homepage (<http://www.aip.org/pub.serve/epaps.html>). **250**

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