To circumvent the limitations of using postmortem brain in molecular assays, we used avidin–biotin binding to couple superparamagnetic iron oxide nanoparticles (SPIONs) (15–20 nm) to phosphorothioate-modified oligodeoxynucleotides (sODNs) with sequence complementary to c-fos and β-actin mRNA (SPION-cfos and SPION-βactin, respectively) (14–22 nm). The Stern–Volmer constant for the complex of SPION and fluorescein isothiocyanate (FITC)-sODN is 3.1 × 10^6 M⁻¹. We studied the feasibility of using the conjugates for in vivo magnetic resonance imaging (MRI) to monitor gene transcription, and demonstrated that these complexes at 40 μg of Fe per kilogram of body weight were retained at least 1 d after intracerebroventricular infusion into the left ventricle of C57Black6 mice. SPION retention measured by MRI as T2* or R2* maps (R2*/H11005) was compared with histology of iron oxide (Prussian blue) and FITC-labeled sODN. We observed significant reduction in magnetic resonance (MR) T2* signal in the right cortex and striatum; retention of SPION-cfos and SPION-βactin positively correlated with c-fos and β-actin mRNA maps obtained from in situ hybridization. Histological examination showed that intracellular iron oxide and FITC-sODN correlated positively with in vivo MR signal reduction. Furthermore, in animals that were administered SPION-cfos and amphetamine (4 mg/kg, i.p.), retention was significantly elevated in the nucleus accumbens, striatum, and medial prefrontal cortex of the forebrain. Control groups that received SPION-cfos and saline or that received a SPION conjugate with a random-sequence probe and amphetamine showed no retention. These results demonstrated that SPION-sODN conjugates can detect active transcriptions of specific mRNA species in living animals with MRI.

Key words: amphetamine; antisense delivery; aptamer; drug addiction; gene transcription; immediate-early genes; nanotechnology; signal transduction

Recently, T₁- and T₂-weighted high-resolution magnetic resonance (MR) techniques have been used to evaluate biological processes. Superparamagnetic iron oxide nanoparticles (SPIONs) (a T₂ contrast agent) can produce significantly strong image contrast by reducing T₂ relaxation time to generate localized signal reduction in T₂-weighted MR images (VanLingen et al., 1998). SPIONs have been shown to be well tolerated as blood-pool contrast agents, and are used in functional magnetic resonance imaging (MRI) studies in live animals, because T₂ signal reduction reflects blood circulation in regions with high cellular activities (Mandeville et al., 1998, 2001). MR contrast agents have also been used to study axonal transport (Petropoulos et al., 1995), and to track transplanted cells in live animals using MRI (Bulte et al., 2002).

Single-stranded phosphorothioate-modified oligodeoxynucleotides (sODNs) with sequences complementary to mRNA have been shown to have specific targeting effects on cerebral mRNAs such as c-fos (Chiasson et al., 1992; Konradi et al., 1994; Liu et al., 1994; Morrow et al., 1999; M. Zhang et al., 1999; Y. Zhang et al., 1999; Tolliver et al., 2000). We have demonstrated sODN uptake in the cerebellum and olfactory bulb of the rat brain after intracerebroventricular infusion (supplemental Fig. SI A, B, available at www.jneurosci.org as supplemental material) (Liu et al., 1994; Cui et al., 1999). To promote the development of ODN-based MRI contrast probes as aptamers for in vivo MR imaging and targeting of cerebral gene transcription, we developed a c-fos probe labeled with the reporter contrast SPION (SPION-cfos) for...
imaging the brains of live C57Black6 mice (see Fig. 1A). We propose that SPION retention reflects gene transcripts in brains of normal animals when reported by SPION-sODN probes with different target sequences (see Fig. 1B). Specifically, this report demonstrates (1) effective linkage between SPION and sODN via conjugation of avidin and biotin, (2) that the stability of SPION-cfos in mouse brain is sufficient for MR assessment, (3) positive correlation of cellular iron oxide and MR image intensity, and (4) that acute amphetamine administration induces specific elevated SPION retention in the forebrains of animals that received SPION-cfos.

**Materials and Methods**

**Contrast probe preparation**

**SPION-NeutrAvidin.** SPION was prepared and purified for these studies in the Athinoula A. Martinos Center for Biomedical Imaging at Massachusetts General Hospital (MGH), as described previously (Shen et al., 1993; Lind et al., 2002). Freshly synthesized SPION was functionalized with cyanogen bromide (Marshall and Rabinowizt, 1976), and linked to NeutrAvidin (NA) in the presence of 1 M sodium cyanoborohydride (both from Pierce Biotechnology, Rockford, IL). The resulting covalently linked products, SPION-NA, was filtered and dialyzed against a 20 X volume of sodium citrate buffer solution (25 mM, pH 8.0), using a Centricon (both from Pierce Biotechnology, Rockford, IL). The resulting covalently linked product, SPION-NA, was stored in an amber-colored bottle at 4°C, at a concentration of 4 mg of iron per milliliter of sodium citrate buffer. Iron concentrations in SPION samples were determined by optical absorbance at 410 nm after treatment with hydrogen peroxide (0.03%) and 6N hydrogen chloride (de Marco et al., 1998).

**Conjugation of biotinylated sODN to SPION-NA.** We synthesized three 5’-biotin-labeled antisense sODN: sODN-cfos, 5’-acgacgtacgtagctgggtttggcaccac-3’ (Liu et al., 1994; Y. Zhang et al., 1999); ODN-Bactin, 5’-acgcagctcagtaacagtcccga-3’ (Schedlich et al., 1997); and sODN-Ran, a randomized sequence, as negative control, 5’-gtagctcagtagctag-3’ (Zhang et al., 2001). All sODN were purified using PAGE. The sequence of sODN-cfos is complementary to rat c-fos mRNA; we used it because the sODN made with mouse sequence contains CCCC, which compromises its stability during storage (our unpublished observation). To directly observe the sODN, we also synthesized sODN with fluorescein isothiocyanate (FITC) on the 5’ terminus and biotin on the 3’ terminus (FITC-sODN-biotin). SPION-NA (250 nmol of Fe) was incubated with the biotinylated sODN (FITC-sODN-biotin) (1 nmol) at room temperature for 30 min and the mixture was filter-dialyzed with three washes of sodium citrate buffer (25 mM, pH 8) in a centrifugal filter device (Microcon YM-30; Millipore). We incubated unconjugated SPION of the same lot from which SPION-sODN had been made with saline. The probes were resuspended in 36 µL of sodium citrate buffer. After conjugation, the probes were stored at 4°C (no longer than 24 h). Immediately before infusion, we added 4 µL of lipofectin (0.1 mg/ml; Invitrogen, San Diego, CA).

**Characterization of SPION-NA and SPION-sODN in vitro.** The core sizes of the conjugates were measured using a Zetasizer particle characterizer system (ZEN3600 with 4 mW 633 He–Ne laser; size range, 0.6 nm to 6 µm; Malvern Instruments, Southborough, MA). The volume fraction (φ) of nanoparticles was used to determine the number (N) of iron atoms per SPION; the volume-weighted diameter of the SPION-cfos nanoparticle in a dilute solution was determined with a modified equation derived by Reynolds et al. (2005). The concentration of SPION was calculated by dividing the iron concentration by the number (N) of iron atoms per nanoparticle.

**The linkage of SPION and sODN using gel shift assay.** The binding capacity for biotin in SPION-NA was determined in triplicate by incubating 100 pmol of FITC-sODN-biotin with 11–171 pmol of SPION-NA at room temperature for 1 h, followed by resolution by electrophoresis in an agarose gel (0.8 or 3%). Because a magnet column cannot partition FITC-sODN-biotin and SPION-cfos, we used gel electrophoresis to separate bound and unbound FITC-sODN-biotin at room temperature. With a secondary absorption peak at 320 nm, FITC-sODN-biotin can be visualized with a UV light source (302/365 nm). Images were captured using a digital camera with a FLSC-400 filter (Alpha Innotech, San Leandro, CA). Binding between FITC-sODN-biotin and SPION-NA will cause the former to migrate more slowly together with SPION and appear as a shift-up in the gel. The appearance of an excess unbound FITC-sODN-biotin band after the saturation point was reached indicated binding capacity; the saturation concentration of FITC-sODN-biotin divided by SPION-NA concentration yielded the number of binding sites (to the nearest whole number). For binding site calculation, please see the example in the supplemental material (available at www.jneurosci.org).

To demonstrate linkage between SPION-NA and FITC-sODN-biotin, we excised four gel slices from each lane, placed each slice in an Eppendorf vial, and then melted and resolubilized the gel to ensure uniform distribution of MR contrast agent. We used a 9.4 tesla MRI system to measure the T2 relaxation times of the gel slices.

**Fluorescence emission and absorption spectra measurements.** Stern–Volmer plots were determined in 25 mM sodium citrate buffer, pH 8.0, by mixing SPION-NA solution (0–6 µM) with free and unbound FITC-sODN-biotin solutions (5 or 10 µM), for solutions with final concentrations ranging from 60 to 900 nm. Fluorescence emission for each solution was measured with a PerkinElmer (Wellesley, MA) Victor 3 V 1420 Multilabel Counter (Wallac) at 450 nm excitation and 510 nm emission. Each concentration of SPION-NA with or without FITC-sODN-biotin was prepared in quadruplicate; two to three concentrations were measured at a time and repeated five times to generate a seven-point curve in the plot. Fluorescence in each concentration was averaged; data were analyzed in GraphPad Prism 4 (GraphPad Software, San Diego, CA). Absorption spectra of FITC-sODN-biotin and SPION-sODN-FITC were measured using a Shimadzu (Tokyo, Japan) UV2101 spectrophotometer; measurements were repeated in triplicate and determined to have two absorption peaks at 495 nm (primary) and 320 nm (secondary).

**Delivery of contrast conjugates**

All procedures and animal care practices adhered strictly to the Association for the Assessment and Accreditation of Laboratory Animal Care, Society for Neuroscience, and institutional guidelines for experimental animal health, safety, and comfort. Male C57Black6 mice (24 g, Taconic, Germantown, NY) were anesthetized with ketamine (42 mg/kg, i.p.) and xylazine (8 mg/kg, i.p.), and the contrast conjugate was delivered to the brain at a dose of 1 µg per mouse (0.04 mg/kg Fe) via intracerebroventricular infusion. The dose administered to each mouse was lower than the nontoxic 1.1 mg/kg Fe dose used in humans (Saini et al., 2000). We used the minimum number of animals necessary, as calculated with a power calculation using the mean and SEM and in-house software by Wayne W. LaMorte (Boston University Medical Center, Boston, MA) (“Sample Size Calculations” available to all animal researchers at MGH).

**Conjugation of biotinylated sODN to SPION-NA.** Unconjugated SPION or SPION-cfos in 2 µL of sodium citrate buffer was delivered (0.2 µL per minute) into the ventricle (stereotactic coordinates from bregma: left–right, −1.0 mm; anterior–posterior, −0.4 mm; dorsal–ventral, −3.0 mm) guided by a stereotactic device (Stoeling, Wood Dale, IL). Afterward, the burr hole in the skull was sealed with bone wax, and the incision was closed with sutures. The infusion dose of SPION was 5 µg per mouse (0.2 mg/kg Fe), because at low dose SPION is not retained in the brain for MR imaging.

To test SPION retention after amphetamine-induced neuronal stimulation, we injected amphetamine (4 mg/kg) or saline (10 ml/kg) intraperitoneally 4 h after SPION-sODN infusion; brain MRI was acquired 3 h later.

**Histology and molecular biological assay**

**Postmortem tissue preparation.** Animals were anesthetized (with ketamine and xylazine as for probe delivery) and the brains were removed from the animals’ skulls and flash-frozen in an optimal cutting temperature compound (Sakura Finetek, Torrance, CA). For MR microscopy, mice were perfused transcardially with 15 ml heparinized saline at a rate of 10 ml/min, and then with 15 ml of freshly prepared paraformaldehyde (PFA) (4%) in 0.1 M PBS, pH 7.4, at a rate of 10 ml/min. Once removed from the animals’ skulls, the brains were stored in PFA solution at 4°C overnight, and then incubated for 16 h in 20% sucrose/PBS solution at 4°C to chase the PFA. Brains were either immediately scanned with high resolution MR microscopy in a 14 tesla MRI system or embedded in paraffin (Cui et al., 1999).
Histological and immunohistochemical staining and molecular assay using in situ hybridization. Postmortem tissue samples were prepared 1 d after infusion, and coronal tissue slices (20–100 mm thick) were prepared from paraffin-embedded blocks as described previously (Liu et al., 1994). A mercury light source, filtered with a 495 nm broad spectrum suitable for wavelengths of 470 nm (excitation) and 525 nm (emission), was used to observe SPION-sODN–FITC. We captured the digital image with a SPOT 1 digital camera (Diagnostic Instruments, Detroit, MI) mounted on an Olympus (Tokyo, Japan) BX-51 microscope (Cui et al., 1999). An adjacent tissue slice was stained with Prussian blue (PB) with 2% potassium ferrocyanide in 2% HCl (Perl’s method) and nuclear fast red (NFR) (Fisher Scientific, Houston, TX) counterstain to detect intracellular iron. For detection of mRNA transcription using in situ hybridization, all protocols including tissue section preparation, 32P-labeled RNA probe (antisense and sense orientation), and autoradiogram exposure were performed as described previously (Cui et al., 1999a).

**MRI protocol**

In vitro MRI of SPION probes. Calibration experiments were performed to investigate the MR T1 and T2 relaxivities (r1 and r2, respectively) of the SPION conjugates used in this study, with different SPION concentrations in 25 mm sodium citrate buffer based phantom fields of strength between 9.4 and 14 tesla, as described previously (Liu et al., 2004). The slopes of the linear fit are the relaxivities (r1, r2) of each contrast probe.

In vivo data acquisition. In vivo image acquisition was performed using a 9.4 tesla MRI scanner (Bruker Avance system, Bruker Biophysics MRI; Bruker, Billerica, MA) at 30 min, 7 h, and 1 d after intracerebroventricular infusion of SPION. Animals were anesthetized with pure O2 and 2% halothane (800 ml/min flow rate) for all time points except that immediately after the intracerebroventricular infusion procedure. We positioned a custom-built 1 cm transmit/receive surface coil on the heads of the animals, which were placed in the prone position in a home-built cradle. Gradient echo (GE) images of constant repetition time (TR) and incremental echo spacing (TE) were acquired at each time point along the axial direction. Acquisition parameters were as follows: TR, 500 ms; TE, 3, 4, 6, 8, and 10 ms; flip angle, 30; 20 0.5 mm slices; 15 15 mm field of view; 128 128 pixel T2* maps were calculated using a pixelwise linear fitting algorithm on the series of images. To compare elevation of mRNA, we constructed R2* maps (the inverse of T2* maps), which showed signal elevation with SPION retention.

**Data analysis.** We selected MR slices from MR data of each animal group (1.7–0.2 mm, relative to bregma) for r test statistical analysis, but avoided those with partial volume artifacts in regions close to the olfactory bulbs and the nose. We used in-house software (Athinoula A. Martinos Center for Biomedical Imaging at MGH) to coregister the images of whole mouse brains, and then superimposed and averaged the R2* maps within designated brain slices. A computer-generated scale is also included in the R2* maps.

**Results**

**Characterization of SPION-sODN**

SPION has an average iron oxide core size of ~10 nm and an average dextran coating thickness of 7 nm, as measured by transmission electron microscopy and light scattering techniques. The design of SPION-sODN is presented in Figure 1A. Table 1 shows both longitudinal and transverse relaxivities (r1 and r2, respectively) for SPION-NA, SPION-cfos (as one example for all SPION-sODNs), and unbound SPION; the SPION-NA and SPION-cfos relaxivities were lower than those of unbound SPION. The average volume-weighted diameter of SPION-cfos was 18 nm at 25°C, and 20 nm at 37°C. Addition of lipofectin to all SPION-sODN before infusion did not significantly change the core size, because <1% measured larger in the presence of lipofectin (Table 1). We calculated the molecular weight of SPION-cfos to be 5.6 107 g per nanoparticles. Each SPION-cfos contains ~6000 iron atoms, based on the relative viscosity of SPION-cfos (1.025 corresponding to 0.4 mg/ml Fe concentration) in sodium citrate buffer at pH 8 (Reynolds et al., 2005). The purity of SPION-NA is presented in the supplemental material (available at www.jneurosci.org). Two sets of evidence on FITC-sODN-biotin binding to SPION-NA are shown here: gel shift assay and Stern–Volmer plot.

**Gel mobility shift on binding**

We presented the electrophoretic migration of one example (SPION-cfos) in agarose gel. Under ordinary light source, SPION appears at 1 and 11 cm, with most of the dye traveling only 1 cm, a distance similar to that traveled by SPION in Figure 2. In lanes in which SPION-NA and FITC-sODN were conjugated and loaded (lanes 2 and 3), fluorescence signal appears at 1 and 11 cm, with most of the dye traveling only 1 cm, a distance similar to that traveled by SPION in Figure 2A, lanes 1–3. This result indicates that there is a band shift typical of DNA–protein binding or conjugation of free sODN and SPION-NA via biotin on sODN and NA on SPION. The appearance of a smear between 1 and 11 cm reflects the size distribution of SPION-NA in Table 1. Variations in fluorescence exist between batches of SPION-NA in lane 2 (batch A, 17 months of age).
and lane 3 (batch B, 2 months of age). The binding capacity remained with long shelf life, although it may show diminished signal intensity (supplemental Fig. S4, available at www.jneurosci.org as supplemental material).

To confirm the presence of SPION in the fluorescent dye of SPION-cfos samples, we measured the T₂ relaxation times in gel slices excised at 0, 1, 2, and 3 cm from the well. Compared with the T₂ relaxation time in the gel without samples (denoted as blank in Fig. 2C), the gel sample from unbound FITC-sODN-biotin (denoted as sODN in Fig. 2C) exhibited no change, indicating SPION was not present. However, Figure 2C shows that the T₂ relaxation times of gels containing SPION-NA were comparatively shortened in gel samples from above the well (0 cm) and 1 cm below the gel; T₂ reduction in the 2 cm gel sample was slightly reduced. The shortened T₂ relaxation times in the gel slice above the wells and at 1–3 cm in the gel containing SPION-sODN in Figure 2C are consistent with fluorescence shift with SPION migration (that is, greater SPION and more T₂ reduction) (Fig. 2A, B). The data demonstrate that SPION-sODN exhibits comigration as a single moiety. We showed binding at equal concentrations for SPION-NA, and FITC-sODN exhibited less fluorescence at 1 cm (Fig. 2B) (also in supplemental Fig. S4, available at www.jneurosci.org as supplemental material), indicating a possible quenching effect when a transition metal is bound to a fluorescence moiety.

Quenching efficiencies
We observed reduction in FITC fluorescence emission when increasing SPION-NA was bound to FITC-sODN-biotin in repeated measurements (Fig. 3A). Using data in Figure 3A, we constructed a Stern–Volmer plot for SPION-sODN to examine the fluorescence quenching effect as a function of iron concentration. Figure 3B demonstrates that the ratio of fluorescence intensity (F₀/F₀) to quencher concentration (iron) detected under various quencher concentrations is linear. The Stern–Volmer constant obtained for iron in SPION was 3.1 × 10⁻³ M⁻¹.

Table 1. Relaxivities (mM⁻¹s⁻¹) of SPION and its conjugates in two different solutions

| 9.4 tesla | 14 tesla | 37°C | 25°C |
|----------|----------|------|------|
| r₁       | r₂       | r₁   | r₂   |
| Volume   | Number   | Volume| Number|
| SPION    | 0.7      | 0.3  | 47   | 21 ± 16 |
| SPION-NA | 0.4      | 0.2  | 25   | 18 ± 11 |
| SPION-cfos| 0.4     | 0.1  | 21   | 20 ± 16 |
| SPION-NA | 0.5      | 0.2  | 29   | 14 ± 4  |
| SPION-cfos| 0.4     | 0.2  | 28   | 15 ± 4  |

The size distribution of the particles was determined using a Zetasizer particle characterization system (Malvern Instruments, MA). SPION was prepared for delivery by three washes in 25 mM citrate buffer using a Millipore filter (30 kDa cutoff). SPION-cfos consists of labeled biotinylated sODN-cfos and SPION-NA. Longitudinal relaxivity is represented by r₁; transverse relaxivity, by r₂. The difference between the r₂ values for SPION and SPION-NA is likely caused by steric hindrance of SPION to surrounding protons due to the presence of NA. SPION-NA has only one binding site for biotinylated sODN-cfos, perhaps because of steric hindrance of SPION.

Colocalization of SPION-NA and s-ODN for MR assessment
To function as a reporter of gene transcription as we envisioned, (1) SPION-sODN must remain linked after delivery to the brain for uptake (Fig. 1A) and (2) SPION retention must be positively correlated with intracellular transcripts and revealed in MR assessment. To test this, we injected either unconjugated SPION or SPION-cfos into the left cerebral ventricle. If the probe complex remains intact, we expect colocalization of sODN and iron oxide 1 d after delivery. The presence of cerebral SPION and sODN were determined by two methods: (1) PB and NFR stains to detect intracellular iron and visualize the nucleus, respectively, and (2) fluorescent microscopy to detect FITC-labeled sODN.

Turnover of SPION in mouse brains
Figure 4 shows MR brain images from live animals obtained at magnetic field strength of 9.4 tesla. Compared with preinfusion brain images (Fig. 4A, baseline), MR signal was reduced from the infusion site across the midline to the right (contralateral) hemisphere immediately (<30 min after SPION-cfos infusion at 1 μg per mouse) (Fig. 4B, top row), distributed at least 4 mm in either direction from the infusion site, and covered the cerebellum and the olfactory ventricle (olfactory bulb), remaining detectable for
Figure 3. Quenching effect of SPION reduces fluorescence emission of the FITC-sODN on SPION. A shows decreasing fluorescence emission of FITC-sODN-biotin on binding to SPION-NA. Error bars indicate SEM. B shows a Stern–Volmer plot constructed for SPION-sODN. $K_q$ is the Stern–Volmer constant calculated from the equation $F_0/F_q = 1 + K_q[Q]$, where $F_0$ and $F_q$ are fluorescence emission at zero ($F_0$) and quencher concentration ($F_q$) of SPION-NA, and $Q$ is the concentration of SPION-NA (Murphy et al., 2004; Li et al., 2005).

at least 1 d after infusion (Fig. 4B, second row). However, relatively little $T_2^*$ signal reduction was observed in animals infused with unconjugated SPION at 1 µg per mouse, except within the infusion track (Fig. 4C). When unconjugated SPION dose was increased to 5 µg; we observed $T_2$ signal reduction at 30 min after infusion (Fig. 4CC, top rows). Unlike SPION-cfos, the majority of unconjugated SPION was no longer present, except in the lateral and fourth ventricles, 1 d after infusion (Fig. 4C, bottom row). SPION was not retained when infused in a mixture of SPION and FITC-sODN-biotin (Fig. 4D), indicating that linkage is important for SPION retention. While unconjugated SPION is rapidly cleared in the cerebral ventricles, SPION-cfos provides an extended window for MR imaging.

Histology: SPION-cfos uptake

Gross examination of PB and NFR stains revealed that Prussian blue staining was stronger in the ipsilateral hippocampus than in the contralateral hippocampus in animals that received SPION-sODN-FITC (Fig. 5, compare A, B). There is weak yet positive signal showing colocalization of sODN and iron in the ipsilateral hippocampus in adjacent postmortem samples (Fig. 5A, C, respectively). At high magnification (Fig. 5A, B, inset panels), we noted particles 1 µm or less in size in the cytoplasm and in regions surrounding the nucleus; cytosolic PB stain was observed as small aggregates (Fig. 5A, B, inset panels, arrows). Figure 5D demonstrates a lack of iron oxide staining in animals that received unconjugated SPION; these animals exhibited no MR signal reduction (Fig. 4C).

Histology: colocalization of iron oxide and SPION-cfos

Figure 5E shows sODN uptake in neurons and their axons (long arrow) between the ventricle and CA neurons (indicated by brackets). We also observed FITC-sODN uptake by cells in the vascular wall (Fig. 5E) and in the CA neuronal formation (Fig. 5G, brackets). Figure 5F shows background fluorescence in the control sample (baseline) of animals that received no infusion. We demonstrate that cells that take up SPION-cfos-FITC are not glia, but neurons (Fig. 5G).

Given the quenching effect of SPION on FITC-sODN, we modified the method by using FITC-Ig against digoxigenin (dig) to detect the presence of sODN with sODN-dig. We infused SPION-cfos-dig (2 µg per mouse) with and without lipofectin and allowed 1 d for uptake. Figure 5, H and I, shows that sODN is present in the cortex and in the CA neuronal formation (green photomicrograph images were converted to black and white with Photoshop software) regardless of whether the infusion included lipofectin. Tissue adjacent to that shown in the inset of Figure 5I (Fig. 5F) also demonstrates the presence of iron oxide in the ventricular wall (asterisk), CA neuronal formation (CA1), dentate gyrus (DG), and vascular wall in the hippocampus. Figure 5K shows PB staining in the cerebellum, including in the Purkinje neurons. The presence of cortical iron oxide has been reported (Liu et al., 2007). The data show that iron oxide and sODN travel at least 1.5 mm (at the hippocampus) to 6 mm (at the cerebellum) from the infusion site.

MR assessment of SPION-cfos correlates with histology of iron oxide

To compare MR assessment to histology with PB stain in Figure 5, A and B, we measured regional $T_2^*$ relaxation time (in milliseconds) at regions of interest (ROIs) in the hippocampus. Compared with the preinfusion baseline hippocampus ($T_2^*$-weighted image shown in Fig. 4Aa), there was no significant difference between regional $T_2^*$ relaxation times (40 ms) of the left and right hippocampus of animals that received unconjugated SPION. In the same ROI of the left (ipsilateral) (LH) and right (contralateral) (RH) hippocampus of animals that received SPION-cfos, regional $T_2^*$ relaxation times were reduced to 7 and 17 ms, respectively (image shown in Fig. 4Bb), suggesting that SPION-cfos is retained in the left hemisphere. Therefore, MR images in Figure 4B correctly reflect regional iron content corresponding to histology of two hemispheres of the same animal brain shown in Figure 5, A and B. The same was observed in three additional brains (data not shown).

SPION-sODN report endogenous gene transcript in live animals

The hippocampus showed the highest uptake within any given brain (Fig. 4B), but we avoided analyzing the hippocampus among different animals for the following reasons. First, given the very small size of the hippocampus and its location in the ventricle, it is very easy to include regions that exhibit ventricular SPION yet to be taken up. Second, the hippocampus contains the hippocampal fissure, where SPION may be detected before its uptake/distribution. We selected the contralateral somatosensory cortices (SSCs) as our ROI for analysis because it is distant from the ventricle and the air–surface interface of the trachea.
(ROI outlines overlay on MR anatomical images). We converted $T_2^*$ relaxation times to $R_2^*$ ($R_2^* = 1/T_2^*$); $R_2^*$ values were used here for statistical analysis of SPION retention. Figure 6A shows statistical analysis of SPION retention in the contralateral SSC of animals that received conjugated and unconjugated SPION probes. Animals that received unconjugated SPION did not exhibit significantly elevated $R_2^*$ values at all time points compared with baseline. SPION retention in animals that received SPION-β-actin and SPION-cfos was significantly higher than baseline as early as 7 h after injection. $R_2^*$ values peaked at 38.5 s$^{-1}$ 1 d after infusion in the contralateral SSC of the animals that received SPION-cfos (Fig. 6A), and returned to baseline values within 2 d (data not shown). We therefore concluded that the clearance time (the time frame spanning the point of infusion to the point at which $R_2^*$ values returned to baseline) was 2 d for the 1 μg dose of SPION-cfos. The $R_2^*$ values in the contralateral SSC of animals that received SPION-β-actin (1 μg) peaked at, and were significantly higher than those of the SPION-cfos group 7 h after infusion, and remained significantly elevated for at least 1 d.

We compared SPION retention to gene transcript maps using in situ hybridization in normal mouse brain samples. The intensity of c-fos mRNA was barely detectable (Fig. 6B) and the mRNA maps of β-actin mRNA were higher than those of c-fos mRNA in normal mouse brains. The intensity of c-fos and β-actin mRNA, shown in Figure 6, B and C, respectively, agrees with SPION retentions of SPION-β-actin and SPION-cfos in the contralateral SSC, 7 h after infusion, as shown in Figure 6A. The data suggest that SPION retention in normal live mice is positively correlated to intracellular mRNA targets of normal mouse brain as hypothesized in Figure 1 B.

**Elevation of SPION retention after acute amphetamine infusion**

We wanted to demonstrate that SPION retention would increase with elevated gene transcripts in those animals treated with targeting SPION-sODN, but not in those infused with nontargeting SPION-Ran (a SPION with sODN of random sequence and therefore no cellular target). We first compared the retention profiles for SPION-cfos and SPION-Ran, both in the same dose, in the striatum of the forebrain (ROI outlines overlay on MR anatomical images), where amphetamine had previously been shown to stimulate expression of c-fos mRNA (Schmauss, 2000; Conversi et al., 2004). SPION retention, measured as $R_2^*$ values, for SPION-Ran in normal brain was within the baseline values (Fig. 6D). SPION-cfos retention was slightly but significantly elevated relative to SPION-Ran 7 h after infusion ($p = 0.004$). These results suggest that the differences in SPION retention between targeting and nontargeting probes were distinguishable using MRI in the contralateral striatum 7 h after intracerebroventricular infusion.

The induction of immediate-early genes by amphetamine is shown using this novel probe and method of detection with MRI. Figure 7 shows $R_2^*$ maps of C57Black6 mice treated with either amphetamine or saline. No elevation in $R_2^*$ was seen in animals that received SPION-Ran, whether with or without acute amphetamine administration (Fig. 7A). Animals that received SPION-cfos and amphetamine exhibited robust elevation of $R_2^*$ in the forebrain (Fig. 7B), particularly in the nucleus accumbens (NAc); $R_2^*$ elevation was less robust in the striatum (CPu) and cingulate and prelimbic/infralimbic cortices [collectively the medial prefrontal cortex (mPFC)], and only little elevation was seen in the SSCs. In the same animals, we observed a gradient of in-
Presence of sODN-dig 1 d after infusion was detected using FITC-IgG against dig (Cui et al., 1999). Previously reported increases in c-ROIs (data not shown). Our observations are consistent with the developments in contrast-enhanced and ligand-targeted MRI complementary to their mRNA targets in autopsy specimens. Recent Assays of gene transcription often use nucleic acid probes complementary to mRNA of neuronal origin (Sagar et al., 1988). The delivery, uptake, and hybridization to mRNA of this ODN in rodent brains has been reported (Liu et al., 1994; Cui et al., 1999; Y. Zhang et al., 1999). When administered at a dose of 1 µg of Fe per mouse via intracerebroventricular injection, SPION-cfos is diffused throughout the brain and selectively retained, as confirmed by recent techniques have made these novel tools for assessing the delivery of biological agents and detecting biological markers in living subjects. DNA-based probe may offer greater variety in aptamer design (Liu et al., 1994; Perez et al., 2002). Among imaging modalities, MRI offers less-invasive image acquisition and enhanced image resolution, and has the potential to become an important tool for assessing delivery of exogenous agents and detecting biological markers in living subjects (Roberts et al., 2000).

We have synthesized several SPION-linked DNA aptamers and demonstrated their application for detecting mRNA in the brains of living animals. Our study focused on one aptamer having sequence complementary to the well characterized c-fos mRNA of neuronal origin (Sagar et al., 1988). The delivery, uptake, and hybridization to mRNA of this ODN in rodent brains have been reported (Liu et al., 1994; Cui et al., 1999; Y. Zhang et al., 1999). When administered at a dose of 1 µg of Fe per mouse via intracerebroventricular injection, SPION-cfos is diffused throughout the brain and selectively retained, as confirmed by increasing R₂* values from the dorsolateral toward ventromedial striata. These results are consistent with distinct patterns of Fos expression in the forebrains of C57Black6 mice associated with acute amphetamine stimulation (Conversi et al., 2004).

Figure 7C demonstrates no significant induction of c-fos expression in the SSC, but a significant elevation in all three ROIs in animals that received SPION-cfos and amphetamine. Because R₂* values in the SSC were not significantly elevated in all treatments, we concluded that our measurements were not significantly influenced by artifacts located outside the brain. Other support comes from control animals that received SPION-Ran, and demonstrated their application for detecting mRNA in the brains of living animals. Our study focused on one aptamer having sequence complementary to the well characterized c-fos mRNA of neuronal origin (Sagar et al., 1988). The delivery, uptake, and hybridization to mRNA of this ODN in rodent brains have been reported (Liu et al., 1994; Cui et al., 1999; Y. Zhang et al., 1999). When administered at a dose of 1 µg of Fe per mouse via intracerebroventricular injection, SPION-cfos is diffused throughout the brain and selectively retained, as confirmed by

Discussion
Assays of gene transcription often use nucleic acid probes complementary to their mRNA targets in autopsy specimens. Recent developments in contrast-enhanced and ligand-targeted MRI techniques have made these novel tools for assessing the delivery of biological agents and detecting biological markers in living subjects. DNA-based probe may offer greater variety in aptamer design (Liu et al., 1994; Perez et al., 2002). Among imaging modalities, MRI offers less-invasive image acquisition and enhanced image resolution, and has the potential to become an important tool for assessing delivery of exogenous agents and detecting biological markers in living subjects (Roberts et al., 2000).
histological assays of iron oxide and immunohistochemical detection of ODN, and corroborated by MRI. SPION unmodified or in mixture with (not linked to) sODN is not retained in the brain, even when given at a dose as high as 5 μg of Fe per mouse; N ≥ 4 each). Multiple cerebral MRIs were acquired as GE MRI in live animals at indicated time points. Preinfusion data serve as baseline (see examples in Fig. 5 for images or saline groups in Fig. 7 for R2* maps). We calculated mean R2* values in an ROI in contralateral SSC to determine SPION retention. No significant differences in R2* values were observed in baseline and animals that received SPION. The p values denote mean R2* values that are significantly different from baseline (*) or significant differences between SPION-cfos and SPION-β-actin (**), calculated using a t test. Based on the mean and SEM from 7 h after infusion, and because power calculation reveals that four animals per group are needed to obtain a statistically significant difference for the effect sizes seen here with 95% confidence and p = 0.01, we measured five mice in each group. Schematic diagrams of anatomic features are within 0.25 mm of each other (see Fig. 6 D). A–J was infused as in Figure 5, except that amphetamine (4 mg/kg body weight) or saline (10 ml/kg) was administered intraperitoneally 4 h later, and MRI was acquired 3 h thereafter. Representative R2* maps (0–200 s) are shown in the scale bar) are from four MR slices (same as Fig. 6). B and C show mRNA maps of c-fos (B) and β-actin (C) gene transcripts in a normal mouse (no infusion) using in situ hybridization and a 32P-labeled antisense RNA probe (Cui et al., 1999a) (N = 4). D shows the profile of SPION retention in the contralateral striatum. Regional SPION retention of SPION-cfos and SPION-Ran (1 μg of Fe per mouse) is shown in the contralateral striatum for ROIs depicted to the right. Error bars indicate SEM.

Figure 6. Temporal profiles of SPION retention in R2* maps. A shows animals infused with SPION-cfos, SPION-β-actin, or unconjugated SPION (1 μg of Fe per mouse; N ≥ 4 each). Multiple cerebral MRIs were acquired as GE MRI in live animals at indicated time points. Preinfusion data serve as baseline (see examples in Fig. 5 for images or saline groups in Fig. 7 for R2* maps). We calculated mean R2* values in an ROI in contralateral SSC to determine SPION retention. No significant differences in R2* values were observed in baseline and animals that received SPION. The p values denote mean R2* values that are significantly different from baseline (*) or significant differences between SPION-cfos and SPION-β-actin (**), calculated using a t test. Based on the mean and SEM from 7 h after infusion, and because power calculation reveals that four animals per group are needed to obtain a statistically significant difference for the effect sizes seen here with 95% confidence and p = 0.01, we measured five mice in each group. Schematic diagrams of anatomic features are within 0.25 mm of each other (see Fig. 6). A–J was infused as in Figure 5, except that amphetamine (4 mg/kg body weight) or saline (10 ml/kg) was administered intraperitoneally 4 h later, and MRI was acquired 3 h thereafter. Representative R2* maps (0–200 s) are shown in the scale bar) are from four MR slices (same as Fig. 6). B and C show mRNA maps of c-fos (B) and β-actin (C) gene transcripts in a normal mouse (no infusion) using in situ hybridization and a 32P-labeled antisense RNA probe (Cui et al., 1999a) (N = 4). D shows the profile of SPION retention in the contralateral striatum. Regional SPION retention of SPION-cfos and SPION-Ran (1 μg of Fe per mouse) is shown in the contralateral striatum for ROIs depicted to the right. Error bars indicate SEM.

Figure 7. Specific elevation of c-fos mRNA in R2* maps after acute amphetamine exposure. SPION-Ran (N = 5 pairs; A) or SPION-cfos (N = 6 pairs; B) was infused as in Figure 5, except that amphetamine (4 mg/kg body weight) or saline (10 ml/kg) was administered intraperitoneally 4 h later, and MRI was acquired 3 h thereafter. Representative R2* maps (0–200 s) are shown in the scale bar) are from four MR slices (same as Fig. 6). C shows profiles of SPION retention in ROIs from R2* maps (see Fig. 6). mPFC, Medial prefrontal cortex; NAc, nucleus accumbens; Cpu, caudate/putamen.
mRNA (Liu et al., 1994) and serves as a primer for specific cDNA reverse transcription (our unpublished observations). Binding of targeting SPION-sODN to its mRNA may create SPION retention (or exclusion delay) by a target-dependent mechanism (Fig. 6). Therefore, the larger aggregates seen under light microscopy with PB staining can be a compartmentalization of iron oxide awaiting exclusion. Compartmentalization could involve endosomes or other cytoplasmic structure (Arbab et al., 2005). This mechanism would explain the elevation of SPION at 7 h for targeting probe (SPION-actin and SPION-cfos) but not nontargeting probe (SPION-Ran).

It has been reported that SPION particles (Combidx; 833 pmol/kg) do not remain in rat brains beyond 3 h (Muldoon et al., 2004), although oligodendroglial cells have been reported to take up an iron oxide-derived MR contrast agent (Bulte et al., 1999). Magnetic nanoparticles covalently attached to DNA were first described by Perez et al. (2002) for detecting DNA–DNA interactions in cellular extracts. We show here a more facile, modular, and universal approach for generating MR reporters for different mRNA targets with DNA–RNA interaction. SPION-cfos and SPION-βactin, at concentrations approximately one-tenth less than that used by Muldoon et al. (2004), are found to be stable and provide an extended window for MR imaging for at least 7–24 h after infusion. Although all types of brain cells are able to take up SODN (Ogawa et al., 1995), we have shown that retention of SPION-cfos and SPION-βactin could be mediated by intracellular mRNA (Fig. 6). These characteristics will be investigated in the future with different probes for different cellular targets.

The intracerebroventricular technique we used to deliver this probe is a convection-enhanced delivery method (Sandberg et al., 2002), and the distribution pattern of SPION-linked ODN delivered by the technique is different from those observed with the use of a viral vector (de Marco et al., 1998; Hommel et al., 2003), in that cellular uptake and hybridization to target mRNA is required for our probe to function as a reporter. The intracranial method of probe delivery results in more uniform distribution with greater reproducibility of uptake and wider reaches into regions away from the infusion site. The presence of SPION in the contralateral hemisphere in animals that received SPION-cfos suggests that distribution of SODN-conjugated SPION is not limited to the infusion site. This delivery method is an improvement over other intracerebral injection techniques (Chiasson et al., 1992; Wahlestedt et al., 1993; Morrow et al., 1999; M. Zhang et al., 1999; Tolliver et al., 2000), as evidenced by the fact that the probes are distributed at least 8 mm anterior–posteriorly (Fig. 4B), and that SPION-cfos migrates to the contralateral hemisphere (Figs. 4B, 5B), possibly by way of the ventricular cerebrospinal fluid, to the perivascular space before axonal transport (Fig. 5E). Therefore, this probe is definitively an improvement over neuronal transport in the guinea pig facial nerve (Petrooulos et al., 1995).

Our delivery method, although invasive for clinical purposes, is appropriate for preclinical neurological assays in drug discovery. Uptake using intracerebroventricular delivery is robust; it is most suitable for applications in preclinical testing, and for use in antisense strategies for gene knock-down, gene therapies, and stem cell tracking. Other methods such as intravenous or intraperitoneal delivery coupled with opening of the blood–brain barrier (BBB) induced by osmotic shock, severe stroke, or trauma, and attaching probes to antibodies against the surface protein of the BBB can presumably be used. However, successful delivery of probes by these methods, although less invasive, is limited to the regions in which the BBB is disrupted by osmotic shock (mammal), or is often disease dependent (stroke and trauma). The intracerebroventricular technique is far more efficient and cost-effective, and can be compared with lumbar puncture in larger animals when proof of concept for the probe is successfully demonstrated by intracerebroventricular delivery and MR imaging in live animals.

Most importantly, we have observed colocalization of sODN and iron oxide in tissue samples taken 1 d after intracerebroventricular infusion; this observation is corroborated by MR signal assessments. We have not been able to show colocalization of SODN-cfos-FITC in the contralateral hemisphere with iron oxide, possibly because of the quenching effect of SPION on FITC, and we are developing a new detection method to overcome this limitation (Fig. 5H). However, iron oxide and sODN-cfos (or SODN-βactin) could be located in the same cells; retention of SPION-cfos and SPION-βactin is mRNA dependent (Fig. 6A, B). Because the half-life of c-fos mRNA, as determined in cell culture, is only ~10 min (Kabnick and Housman, 1988), the enhanced stability of SPION retention at 7–24 h in the brains of animals that received SPION-cfos (or SPION-βactin), but not unconjugated SPION or SPION-Ran, suggests the extended window of MR imaging may be a result of heteroduplex formation between mRNA and targeting SPION-sODN. This is supported by the fact that SODN-cfos binds to c-fos mRNA both in vitro and in vivo, especially in the neuronal formation where FOS antigens are present in abundance (Liu et al., 1994). Furthermore, SPION-cfos forms a hybrid with c-fos mRNA in mouse brains (our unpublished observations). Lastly, elevated SPION retention in the forebrains of animals that received SPION-cfos, but not those that received SPION-Ran after acute administration of amphetamine corroborates elevation of c-fos mRNA.

Amphetamine alters the expression of several genes. The exact action of amphetamine is not totally understood, but it stimulates brain cells through several membrane receptors in the striatum and induces signal transduction by phosphorylation of cAMP response element binding protein (CREB). CREB activates the transcription of several genes, one of which is c-fos (Konradi et al., 1994; Hyman and Malenka, 2001). We also detected mRNA elevations in MRI using SPION-sODN probes in additional disease models (our unpublished observations). In the current study, the elevation of c-fos mRNA by amphetamine increases the hybridization with SPION-cfos probe and R* values in regions that show neuronal activation (Fig. 7). In conclusion, we designed target-specific RNA sensors and demonstrated their usefulness for in vivo monitoring of gene transcripts by time-lapse imaging of live animals.

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