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On-chip manipulation of nonmagnetic particles in paramagnetic solutions using embedded permanent magnets

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Abstract This study develops a method for embedding permanent magnets into poly(dimethylsiloxane) (PDMS)based microfluidic chips. Magnets can be brought very close to the planar microchannels for enhanced magnetic field and field gradients, which enables on-chip continuous-flow manipulation of nonmagnetic particles in typical paramagnetic solutions. We performed a systematic study of the transport of polystyrene particles suspended in manganese (II) chloride (MnCl₂) solutions through a rectangular microchannel. Owing to their smaller magnetization than the suspending fluid, particles experience negative magnetophoresis and are deflected away from the magnet. The effects of particle position (relative to the magnet), particle size, MnCl₂ salt concentration, and fluid flow velocity on the horizontal magnetophoretic deflection are examined using a combined experimental and theoretical approach. The experimental results agree quantitatively with the predictions of an analytical model. The demonstrated nonmagnetic particle deflection may be used with the potential to focus and sort cells in lab-on-a-chip for bio-applications.

Keywords Microfluidics · Magnetophoresis · Nonmagnetic particle · Paramagnetic solution · Lab on a chip

1 Introduction

Magnetophoresis is a simple, efficient, and low-cost technique for particle manipulation in microfluidic devices

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(Pamme 2006; Gijs et al. 2010). It refers to the induced motion of particles in a non-uniform magnetic field (Jones 1995). Analogous to dielectrophoresis (Pohl 1978), magnetophoresis can be either positive or negative depending on whether the particle is more or less magnetizable than the suspending medium (Kirby 2010). Magnetic particles suspended in nonmagnetic solutions experience positive magnetophoresis and are directed along the magnetic field gradient toward a magnet (Pamme and Manz 2004; Gijs 2004; Pamme and Wilhelm 2006). On the contrary, nonmagnetic particles suspended in magnetic solutions undertake negative magnetophoresis and are pushed away from a magnet (Rosensweig 1985; Friedman and Yellen 2005). The former phenomenon has been widely employed to isolate and sort cells or biomolecules in microfluidic devices by selectively labeling the target ones with functionalized magnetic beads (Liu et al. 2009; Ando et al. 2010) or by utilizing their intrinsic paramagnetic feature (Han and Frazier 2006; Jung et al. 2010).

Ferrofluids (Rosensweig 1987; Erb and Yellen 2009) and paramagnetic salts (Suwa and Watarai 2011) are the magnetic solutions frequently employed to manipulate nonmagnetic particles that cover the majority of the synthetic and biological particles. As no magnetic tagging is needed, this negative magnetophoresis-based method can greatly facilitate the sample preparation and post-analysis, both of which are critical steps in lab on a chip for bioapplications. Ferrofluids have been demonstrated to pattern (Yellen et al. 2003, 2005; Yellen and Friedman 2004; Halverson et al. 2006), assemble (Feinstein and Prentiss 2006; Erb et al. 2009; Krebs et al. 2009; Li and Yellen 2010), concentrate (Fateen 2002; Erb and Yellen 2008; Annavarapu 2010), deflect (Sharpe 2004; Zhu et al. 2011), and separate (Kose et al. 2009; Zhu et al. 2010) particles with various sizes. They are, however, opaque liquids, and



so fluorescent staining is usually necessary in order to visualize the suspending particles.

In contrast, paramagnetic salts such as manganese(II) chloride (MnCl₂) and gadolinium(III) chloride (GdCl₃) solutions are transparent, which is beneficial to particle handling. However, their magnetic susceptibility is usually much smaller than that of ferrofluids (Mirica et al. 2009). Therefore, the salt concentration must be kept high in order to generate decent particle magnetophoresis, which makes the paramagnetic solution non-biocompatible (Rodriguez-Villarreal et al. 2011). Alternatively, strong magnet(s) [e.g., a superconducting magnet (Tarn et al. 2009)] must be employed to provide large magnetic fields. Another way is to bring magnet(s) very close to the microchannel that contains the particle suspension or even in contact with the suspension itself (Winkleman et al. 2004), so that the large magnetic field gradients can be achieved. Pamme's group (Peyman et al. 2009; Tarn et al. 2009; Rodriguez-Villarreal et al. 2011) and Watarai's group (Watarai and Namba 2001, 2002; Watarai et al. 2004) have each demonstrated continuous-flow control of particles and cells in paramagnetic solutions by the use of two closely facing magnets around a capillary. In addition, Park's group (Kang et al. 2008; Hahn and Park 2011) used an on-chip Nickel microstructure near the microchannel to locally enhance both the magnetic field and the field gradients for particle separation in paramagnetic buffers.

We develop herein a method to place permanent magnets to within 300-µm distance of an on-chip planar microchannel by embedding them into poly(dimethylsiloxane) (PDMS) directly. This enables the continuous-flow manipulation of nonmagnetic particles in typical paramagnetic solutions. To demonstrate this, we investigate the negative magnetophoretic deflection of polystyrene particles in MnCl₂ solutions through a rectangular microchannel. We also develop an analytical model to understand the observed particle motion, which is validated by the acquired experimental results. This model is believed to be applicable to previous studies on particle magnetophoresis in paramagnetic solutions.

2 Experiment

2.1 Device fabrication

Figure 1 shows a picture of the microfluidic device used in our experiment. The straight microchannel (filled with a green food dye for clarity) is 2-cm long, 200-µm wide and 25-µm deep. It was fabricated using a modified soft lithography method in order to embed permanent magnets into PDMS. Prior to the dispense of liquid PDMS over the

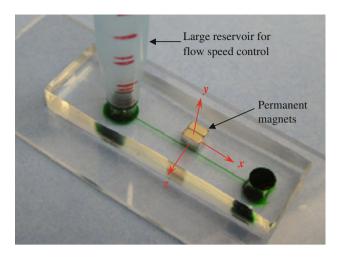


Fig. 1 Picture of the microfluidic chip with permanent magnets embedded in PDMS. The coordinate system, based on which the magnetic field and field gradients are determined, originates from the center of the permanent magnet whose magnetization direction is in line with the z coordinate

channel master (Zhu et al. 2009), two stacked neodymium (NdFeB) permanent magnets (B221, K&J Magnetics, Inc.) were positioned 260 µm away from the edge of the microchannel. They were fixed to the substrate of the master by another permanent magnet (B421, K&J Magnetics, Inc.) from underneath the petri dish that held the master. These steps were performed under the microscope for a precise control. After the regular curing step, the magnet underneath the petri dish was removed, and the cured PDMS was carved out and bonded to a glass slide forming the microfluidic chip. A post-bonding baking was found to enhance the sealing quality of the microchannel. The embedded magnets each has a dimension of $1/8'' \times 1/8''$ $8'' \times 1/16''$ (thick), and is magnetized through the thickness (i.e., z direction) that is perpendicular to the flow direction as seen in Fig. 1.

2.2 Preparation of particle suspensions

Manganese (II) chloride (MnCl₂) crystal was purchased from Fisher Scientific. It was dissolved in pure water to a concentration of 2 M, which was then stored in refrigerator for later uses. Prior to experiments, the stored MnCl₂ solution was diluted to 1 M, 200 mM, or 40 mM using pure water. Polystyrene particles of 5, 10, and 15 μ m in diameters were obtained from Sigma-Aldrich. They are all originally packed as 10% solids in water with the size non-uniformity being less than 2%. Each type of these non-magnetic microparticles was re-suspended in a MnCl₂ solution (at one of the above-mentioned concentrations) to a final number density of about 1 \times 10⁶ particles/ml.



2.3 Particle manipulation and visualization

The particle suspensions in MnCl₂ solutions were driven through the microchannel by the liquid height difference in the upstream and downstream reservoirs. During experiment, the solution in the downstream reservoir was vacated. The liquid height in the upstream reservoir was varied to achieve different flow speeds, which were first estimated through theoretical calculations and then verified via experimental tracking of individual particles. Particle motion was visualized and recorded using an inverted microscope (Nikon Eclipse TE2000U, Nikon Instruments, Lewisville, TX) equipped with a CCD camera (Nikon DS-Qi1Mc). The obtained images were then processed using the Nikon imaging software (NIS-Elements AR 2.30).

3 Theory

3.1 Magnetic field-induced particle deflection

The magnetic force, \mathbf{F}_{m} , on a particle (either magnetic or nonmagnetic) induced by magnetic field gradients is given by (Jones 1995; Pamme 2006; Suwa and Watarai 2011):

$$\mathbf{F}_{\mathrm{m}} = 3\mu_{0}V_{\mathrm{p}}\frac{\chi_{\mathrm{p}} - \chi_{\mathrm{f}}}{3 + \chi_{\mathrm{p}} + 2\chi_{\mathrm{f}}}(\mathbf{H} \bullet \nabla)\mathbf{H}$$
$$\cong \mu_{0}V_{\mathrm{p}}(\chi_{\mathrm{p}} - \chi_{\mathrm{f}})(\mathbf{H} \bullet \nabla)\mathbf{H} \tag{1}$$

where $\mu_0 = 4\pi \times 10^{-7}$ H/m is the permeability of free space, V_p is the volume of the particle, χ_p is the magnetic susceptibility of the particle, χ_f is the magnetic susceptibility of the suspending fluid, and H is the magnetic field at the particle center. The formula after the second equal sign is valid with negligible errors because both χ_p and χ_f are at least three orders of magnitude smaller than 1 for the nonmagnetic particles and paramagnetic solutions used in this study. Moreover, as $\chi_p < \chi_f$, nonmagnetic particles suspended in paramagnetic solutions are directed by $\boldsymbol{F}_{\boldsymbol{m}}$ against the magnetic field gradient, i.e., along the direction of decreasing magnetic field magnitude. For a rectangular permanent magnet with magnetization along the z direction, the analytical formulae for the three components of **H**, i.e., H_x , H_y and H_z , are given by (Furlani 2001):

$$H_{x}(x,y,z) = \frac{M_{s}}{4\pi} \sum_{i=1}^{2} \sum_{j=1}^{2} (-1)^{i+j} \ln \left\{ \frac{(y-y_{1}) + \left[(x-x_{i})^{2} + (y-y_{1})^{2} + \left(z-z_{j} \right)^{2} \right]^{1/2}}{(y-y_{2}) + \left[(x-x_{i})^{2} + (y-y_{2})^{2} + \left(z-z_{j} \right)^{2} \right]^{1/2}} \right\}$$
(2)

$$H_{y}(x, y, z) = \frac{M_{s}}{4\pi} \sum_{i=1}^{2} \sum_{j=1}^{2} (-1)^{i+j} \ln \left\{ \frac{(x - x_{1}) + \left[(x - x_{1})^{2} + (y - y_{i})^{2} + (z - z_{j})^{2} \right]^{1/2}}{(x - x_{2}) + \left[(x - x_{2})^{2} + (y - y_{i})^{2} + (z - z_{j})^{2} \right]^{1/2}} \right\}$$

$$(3)$$

$$H_{z}(x, y, z) = \frac{M_{s}}{4\pi} \sum_{i=1}^{2} \sum_{j=1}^{2} \sum_{k=1}^{2} (-1)^{i+j+k} \tan^{-1} \times \left\{ \frac{(x - x_{i})(y - y_{j})}{(z - z_{k}) \left[(x - x_{i})^{2} + (y - y_{j})^{2} + (z - z_{k})^{2} \right]^{1/2}} \right\}$$
(4)

where the coordinate system originates from the center of the magnet, and $M_{\rm s}=1.05\times 10^6$ A/m is the residual magnetization of the permanent magnet as calculated from the residual magnetic flux density, $B_{\rm s}$ (=1.32 T as per the manufacturer), through $M_{\rm s}=B_{\rm s}/\mu_0$. Other symbols involved in the magnetic field equations are $x_1=x_{\rm m}$, $x_2=-x_{\rm m}$, $y_1=y_{\rm m},y_2=-y_{\rm m},z_1=z_{\rm m}$, and $z_2=-z_{\rm m}$ with $x_{\rm m},y_{\rm m}$, and $z_{\rm m}$ being one half of the dimensions of the magnet in the x,y, and z directions, respectively.

The total magnetic field of the two stacked permanent magnets in Fig. 1 is simply the superposition of **H** of each magnet given in Eqs. 2-4. Fig. 2 shows the contours of the total magnetic field magnitude in the horizontal (at $y = -y_m + h_c/2$ with h_c being the microchannel height, left plot) and vertical (at x = 0, right plot) planes of the microchannel. Table 1 summarizes the parameters involved in the calculation. It is evident that the magnets generate strong magnetic field gradients nearby in all the three directions. Moreover, the further away from the center of the magnets, the smaller the local magnetic field is. Therefore, nonmagnetic particles should be deviated from the fluid flow in the x-direction, and be deflected

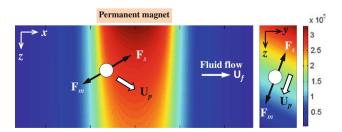


Fig. 2 Force analyses on a nonmagnetic particle suspended in $MnCl_2$ solution in the horizontal (*left plot*) and vertical (*right plot*) planes of a microchannel. The background shows the contours of magnetic field strength (in the unit of H/m) in the absence of the nonmagnetic particle. \mathbf{F}_m and \mathbf{F}_s denote the magnetic and Stokes drag forces, respectively. \mathbf{U}_f and \mathbf{U}_p denote the fluid and particle velocities, respectively. The microchannel and magnet are not drawn to scale

Table 1 List of the parameters used in the an	dytical model
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Parameter		Description	Value	Unit
Magnet	$M_{ m s}$	Residual magnetization	1.05×10^{6}	A/m
	$2x_{\rm m}$	Length	3.176	mm
	$2y_{\rm m}$	Height	3.176	mm
	$2z_{\rm m}$	Thickness (polar direction)	1.588	mm
		Magnets-channel distance	260	μm
MnCl ₂ solution	c	Molar concentration	40, 200, 1000	mM
	$ ho_{ m f}$	Mass density	1050	kg/m ³
	$\eta_{ m f}$	Dynamic viscosity	1×10^{-3}	kg/m/s
	χ_{f}	Magnetic susceptibility	Eq. (9)	
Nonmagnetic particles	d	Particle diameter	5, 10, 15	μm
	$\chi_{ m p}$	Magnetic susceptibility	-5×10^{-6}	
	$l_{\rm c}$	Channel length	2	cm
Microchannel	$w_{\rm c}$	Channel width	200	μm
	h_{c}	Channel height	25	μm
	$U_{ m avg}$	Average flow speed	200, 400, 600	μm/s

along both the positive z and negative y directions as illustrated by the force analyses in Fig. 2. The result is a three-dimensionally focused particle stream flowing near the outer bottom corner of the microchannel. However, since the depth (25 μ m) of the current channel is much smaller than its width (200 μ m), particles are expected to be deflected toward the bottom wall much more quickly than to the outer sidewall. Therefore, for simplicity, we assume in our model (see Sect. 3.2 for detail) that particles are brought into touch with the bottom wall momentarily by the vertical magnetic force and then move through the magnets region only in the horizontal plane.

The particle *deflection* in the horizontal plane (i.e., x–z plane) of the microchannel is determined by the ratio of the particle velocities, U_p , perpendicular and parallel to the flow

Deflection =
$$\frac{U_{p,z}}{U_{p,x}} = \frac{U_{m,z}}{U_f + U_{m,x}}$$
 (5)

where $U_{\rm f}$ is the axial flow velocity (White 1991)

with U_{avg} being its cross-sectional area-average

$$U_{\text{avg}} = \frac{\rho_{\text{f}}g\Delta h}{l_{\text{c}}} \frac{8h_{\text{c}}^2}{\eta_{\text{f}}\pi^4} \left\{ \left[1 - \frac{2h_{\text{c}}}{\pi w_{\text{c}}} \tanh\left(\frac{\pi w_{\text{c}}}{2h_{\text{c}}}\right) \right] - \frac{1}{81} \left[1 - \frac{2h_{\text{c}}}{3\pi w_{\text{c}}} \tanh\left(\frac{3\pi w_{\text{c}}}{2h_{\text{c}}}\right) \right] \right\}$$
(7)

In the last two equations, ρ_f is the density of the MnCl₂ solution, g is the gravitational acceleration, Δh is the liquid height difference in the upstream and downstream reservoirs of the microchannel that provides the pressuredrop for driving the particle suspension, l_c , h_c , and w_c are the length, height, and width of the microchannel, and η_f the dynamic viscosity of the suspending solution. The auxiliary coordinates y' and z' in Eq. 6 originate from the center of the channel cross-section and are parallel to the y and z coordinates for the magnet (see Fig. 1), respectively.

The magnetophoretic particle velocity, \mathbf{U}_{m} , in Eq. 5 can be obtained by balancing the magnetic force, \mathbf{F}_{m} , in Eq. 1 with the Stokes drag force (see \mathbf{F}_{s} in Fig. 2),

$$U_{\rm f} = \frac{\pi U_{\rm avg}}{2} \left\{ \left[1 - \frac{\cosh\left(\frac{\pi c'}{h_{\rm c}}\right)}{\cosh\left(\frac{\pi w_{\rm c}}{2h_{\rm c}}\right)} \right] \cos\left(\frac{\pi y'}{h_{\rm c}}\right) - \frac{1}{27} \left[1 - \frac{\cosh\left(\frac{3\pi z'}{h_{\rm c}}\right)}{\cosh\left(\frac{3\pi w_{\rm c}}{2h_{\rm c}}\right)} \right] \cos\left(\frac{3\pi y'}{h_{\rm c}}\right) \right\}$$

$$\left\{ \left[1 - \frac{2h_{\rm c}}{\pi w_{\rm c}} \tanh\left(\frac{\pi w_{\rm c}}{2h_{\rm c}}\right) \right] - \frac{1}{81} \left[1 - \frac{2h_{\rm c}}{3\pi w_{\rm c}} \tanh\left(\frac{3\pi w_{\rm c}}{2h_{\rm c}}\right) \right] \right\}^{-1}$$

$$(6)$$



$$\mathbf{U}_{\mathrm{m}} = \frac{\mathbf{F}_{\mathrm{m}}}{3\pi\eta_{\mathrm{f}}df_{\mathrm{D}}} = \frac{\mu_{0}d^{2}(\chi_{\mathrm{p}} - \chi_{\mathrm{f}})(\mathbf{H} \bullet \nabla)\mathbf{H}}{18\eta_{\mathrm{f}}f_{\mathrm{D}}}$$
(8)

where d is the particle diameter, and $f_{\rm D}=3.1$ is the drag coefficient accounting for the channel wall's retardation effect on particle motion (Happel and Brenner 1973). The magnetic susceptibility of the MnCl₂ solution is assumed to increase linearly with the concentration of MnCl₂, c, via (Mirica et al. 2009).

$$\gamma_f = 1.877 \times 10^{-4} c - 9 \times 10^{-6} \tag{9}$$

Therefore, \mathbf{U}_{m} and hence the particle *deflection* should increase with increasing particle size and MnCl₂ concentration. In addition, lowering the flow velocity should also enhance the particle deflection according to Eq. 5.

3.2 Simulation of particle trajectory

We developed an analytical model to simulate the trajectory of nonmagnetic particles in paramagnetic solutions in response to magnetic field gradients. As explained earlier, this model assumes that particles are in touch with the bottom wall of the microchannel and travel only in the horizontal plane (i.e., x–z plane) for simplicity. The instantaneous position of a particle, \mathbf{r}_p , was obtained by integrating the particle velocity over time,

$$\mathbf{r}_{p} = \mathbf{r}_{0} + \int_{0}^{t} \left[\mathbf{U}_{f}(t') + \mathbf{U}_{m}(t') \right] dt'$$
 (10)

where \mathbf{r}_0 is the initial location of the particle, and t is the time coordinate. It is noted that both the fluid velocity, U_f , and the magnetophoretic particle velocity, U_m , are dependent on position, and so vary with time during the particle migration. Inertia is neglected in Eq. 10 as both the calculated fluid and particle Reynolds numbers are much smaller than 1 in the experimental conditions (Di Carlo et al. 2007). A custom-written Matlab® program was employed to determine the particle position with respect to time and to plot the particle trajectory. A total of twenty evenly distributed points over the width were picked at the entrance of the microchannel as the initial particle positions. The magnetic field components were calculated from Eqs. 2-4 by setting $y = -y_m + d/2$, i.e., the y-coordinate value of the particle center. This same y value was also assigned to Eq. 6 to determine the axial fluid velocity $U_{\rm f}$. All the parameters involved in the model are listed in Table 1 unless otherwise stated in the text.

4 Results and discussion

4.1 Evolution of horizontal particle deflection

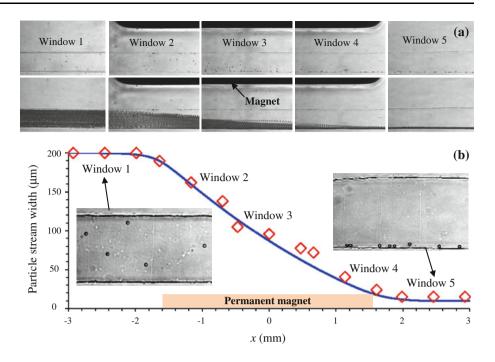
To understand how the magnetic deflection evolves when particles approach and move by the permanent magnets, we studied 10 µm particle motions in 1 M MnCl₂ solution through a row of five observation windows along the channel length. The relative positions of these windows with respect to the magnets are 0.5 mm upstream (Window 1), align with the front edge (Window 2), center (Window 3), align with the back edge (Window 4), and 0.5 mm downstream (Window 5). Figure 3a shows the top-view images (top row for snapshot, bottom row for superimposed) in the five observation windows at an average flow speed of $U_{\rm avg} = 800 \ \mu \text{m/s}$. One can see that particles follow the fluid flow in Window 1, indicating negligible magnetic force when particles are 0.5 mm ahead of the magnets. When they approach the front edge of the magnets in Window 2, particles start being deviated from the axial flow direction and acquire a noticeable deflection away from the magnets. This horizontal deflection grows continuously to nearly the entire channel width as particles move through the magnets region, which is clear from the images in Windows 2-4. It vanishes when particles move into Window 5. Therefore, the effective region for particle magnetophoresis is restricted to only the part of the microchannel that is right beside the magnets.

The observed lengthwise evolution of particle deflection in Fig. 3a is accurately predicted by the analytical model as demonstrated in Fig. 3b, where the experimentally measured (symbols, determined from the superimposed images) and theoretically predicted (curve) widths of the focused particle stream are quantitatively compared. The experimental error in measuring the stream width is less than 10 µm, which is too small to be clearly illustrated in Fig. 3b and thus, not included. A good agreement is observed for the results in all the five observation windows. Moreover, the inset images in Fig. 3b demonstrate that in Window 1 particles appear in different brightness and thus should stay in different focal planes, i.e., different vertical planes of the microchannel. In contrast, all the particles in Window 5 exhibit a nearly identical appearance and, are thus, within the same focal plane. This indicates a threedimensional focusing effect due to the horizontal and vertical magnetophoretic deflections as noted in the Theory section (see also Fig. 2).

The observation in Fig. 3 is also consistent with the axial distribution of the predicted magnetophoretic particle velocity, $\mathbf{U}_{\rm m}$, along the channel centerline. As seen in Fig. 4, the cross-stream component, $U_{\rm mz}$, acquires a positive value (on the order of 10 μ m/s) only within



Fig. 3 Lengthwise evolution of the magnetophoretic deflection of 10 µm polystyrene particles suspended in 1 M MnCl₂ solution at an average flow speed of 800 µm/s: snapshot (top row) and superimposed (bottom row) images in the five consecutive observation windows along the channel length (a); comparison of the experimentally measured (symbols) and theoretically predicted (curve) widths of the particle stream along the flow direction (b). The insets of (b) show the magnified images of particles in Windows 1 and 5, respectively. The position of the two stacked permanent magnets is indicated in the superimposed image in Window 3 of (a) and is also highlighted in (b)



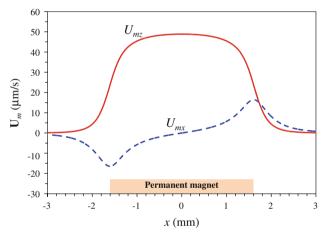


Fig. 4 Axial distribution of the theoretically predicted magnetophoretic particle velocity, $U_{\rm m}$, along the centerline of the microchannel. All the conditions are referred to Fig. 3

0.5 mm distance before and after the magnet (i.e., -2 mm < x < +2 mm), which explains the question of why no apparent particle deflection is observed in Windows 1 and 5 of Fig. 3. The streamwise magnetophoretic velocity, $U_{\rm mx}$, also obtains a non-zero value in the same range of x as $U_{\rm mz}$ though at a smaller magnitude. Moreover, $U_{\rm mx}$ varies from negative in the upstream half of the magnet, which hinders the particle motion, to positive in the downstream half, which propels the particle.

4.2 Particle size effect on particle deflection

Equation 8 indicates that the magnetophoretic velocity is proportional to particle diameter squared. As a result, the

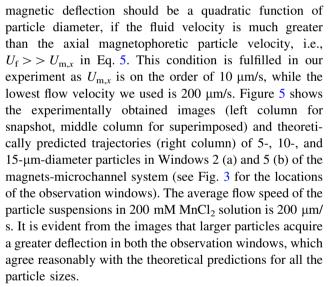
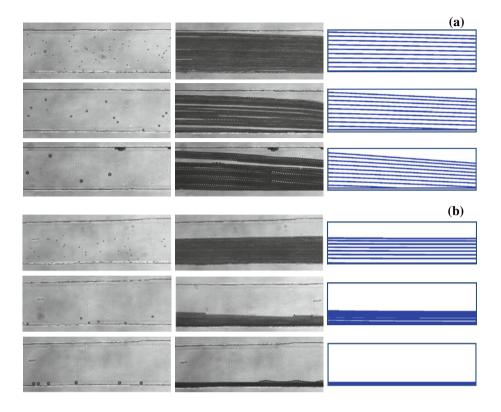
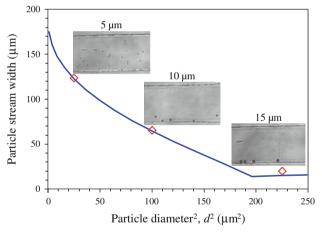


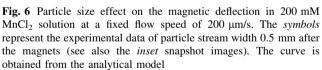
Figure 6 shows a quantitative comparison of the experimentally measured (symbols, see also the inset images) and theoretically predicted (curved) stream widths of different particles 0.5 mm after the magnets (i.e., in observation Window 5). It is noted that the horizontal axis is the particle diameter squared. The agreement is good for all the three particle sizes. Moreover, the stream widths of 5 and 10 μ m particles indeed approximately scale with the square of their diameters. The deviation from the second order dependence for particles smaller than 5 μ m particles is probably attributed to the variation of the magnetic field and fluid velocity at the particle center with particle size. For particles larger than 14 μ m, the model predicts a full-width deflection and so the particle stream width becomes essentially equal to the particle diameter. This prediction is



Fig. 5 Illustration of particle size effect on the deflection of nonmagnetic particles in Window 2 (a) and Window 5 (b) of the magnetsmicrochannel system (see Fig. 3 for the locations of the observation windows), where the top, middle, and bottom rows in each panel are for particles of 5, 10, and 15 µm in diameter, respectively. The columns from left to right show the snapshot image, superimposed image, and theoretically predicted trajectories of the particles. respectively. All the particles were re-suspended in 200 mM MnCl₂ solution. The average flow speed was fixed at 200 µm/s







evidenced from the inset image in Fig. 6, where 15 μ m particles all line the channel sidewall as a consequence of the negative magnetophoretic deflection.

4.3 MnCl₂ concentration effect on particle deflection

The molar concentration of MnCl₂ salt, c, affects the magnetic susceptibility, χ_f , density, ρ_f , and viscosity, η_f , of the solution, among which the changes in the latter two

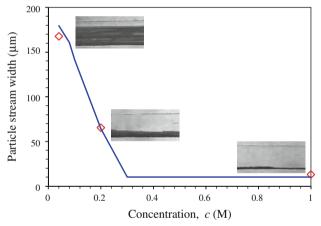


Fig. 7 MnCl₂ concentration effect on the deflection of 10 μ m nonmagnetic particles at an average flow speed of 200 μ m/s. The *symbols* represent the experimental data of particle stream width measured from the corresponding superimposed images. The curve is obtained from the analytical model

properties are much smaller than that in χ_f . Therefore, we neglected the MnCl₂ concentration effects on the fluid density and viscosity in the analytical model. The magnetophoretic particle deflection is expected to be linearly proportional to c. This analysis is verified in Fig. 7, where the experimentally measured stream widths of 10 μ m particles suspended in 40 mM, 200 mM, and 1 M MnCl₂ solutions (symbols, see also the inset superimposed images) closely match the theoretical prediction (curve). The



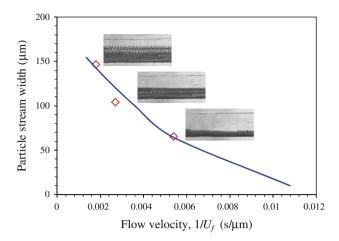


Fig. 8 Flow velocity effect on the deflection of 10 μm nonmagnetic particles suspended in 200 mM MnCl₂ solution. The *symbols* represent the experimental data of particle stream width measured from the corresponding superimposed images. The curve is obtained from the analytical model

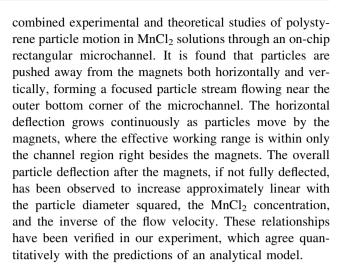
model predicts that $10~\mu m$ particles should achieve a full-width deflection when the $MnCl_2$ concentration is larger than 300~mM, which is supported by the experimental data at 1~M concentration. Below 300~mM, the particle stream width decreases (equivalently, particle deflection increases) linearly with the $MnCl_2$ concentration as expected.

4.4 Flow effect on particle deflection

Equation 5 indicates that particle deflection should decrease linearly with increasing flow speed of the MnCl₂ solution as long as $U_{\rm f} >> U_{\rm m,x}$. To test this, we studied the deflection of 10 µm particles in 200 mM MnCl₂ solution when the average flow speed was varied from 200 to 600 µm/s. The experimental results (symbols and inset images) of the particle stream width are compared with the theoretical prediction (curve) in Fig. 8, where a reasonable agreement is found for the three tested flow velocities. It is noted that the horizontal axis is the inverse of the flow velocity. We see that to a good approximation, the predicted particle stream width is inversely proportional to the flow velocity, which is consistent with the experimental results.

5 Conclusion

We have developed a method to embed permanent magnets into PDMS, and demonstrated that the magnet-microchannel distance can be precisely controlled down to $260~\mu m$. This enables a continuous-flow manipulation of nonmagnetic microparticles in paramagnetic solutions on PDMS-based microfluidic chips. We have performed



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