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Isolation of DNA Aptamers Using Micro Free Flow Electrophoresis

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Abstract

A micro free flow electrophoresis (μ FFE) device was used to select DNA aptamers for human immunoglobulin E (IgE). The continuous nature of μ FFE allowed 1.8×10^{14} sequences to be introduced over a period of 30 minutes, a 300-fold improvement in library size over capillary electrophoresis based selections (CE-SELEX). Four rounds of selection were performed within four days. Aptamers with low nM dissociation constants for IgE were identified after a single round of μ FFE selection.

Introduction

Aptamers are single stranded DNA or RNA molecules that fold into unique structures that promote selective, high-affinity interactions with specific target molecules.¹ Aptamers are isolated using an *in vitro* process commonly referred to as Systematic Evolution of Ligands by EXponential enrichment (SELEX).^{2, 3} Generally, the SELEX process consists of iterative rounds alternating between affinity enrichment of binding sequences and PCR amplification. Typically, aptamers with low nM to high pM dissociation constants are isolated after 10–15 rounds of selection. Aptamers have been selected for a variety of targets, including cells,^{4–6} proteins,^{7, 8} antibiotics,^{9–11} and small molecules such as amino acids^{12–14} and biological cofactors.^{15, 16} Aptamer applications range from purification,^{17–19} detection,^{20–23} and quantification^{22, 24, 25} of targets in basic research to diagnostic agents^{26, 27} and drug candidates^{28–32} in clinical applications.

Conventionally, affinity selections in SELEX are performed using nitrocellulose membranes or affinity chromatography, requiring manual manipulation of relatively large sample volumes and providing ample opportunity for non-specific interactions with solid surfaces. Typically 10–15 selection cycles are required, making the SELEX process labour intensive and time consuming. Since the earliest aptamer publications,^{2, 3} great efforts have been made to develop protocols that minimize solution volumes, are simple and fast, and are compatible with automated processes. Cox *et al.* were the first to automate the SELEX process by incorporating the separation, PCR and purification apparatus into a robotic pipetting worksurface.^{33, 34} Twelve cycles of selection were completed within 42 hours, reducing the selection time from several weeks to less than two days.

Recent developments in microfluidics provide further opportunities for enhancing the SELEX process. It is anticipated that a high through-put lab-on-a-chip device that combines

high separation efficiency, increased PCR speed, low sample volume, minimal contamination, and automation will be possible.³⁵⁻³⁸ Towards this goal several microfluidic devices have been reported for isolating aptamers. A microfluidic device patterned with sol-gel droplets on individual microheaters was developed.³⁹ This device can selectively generate aptamers for multiple targets simultaneously within the sol-gel droplets, significantly improving throughput. A continuous-flow magnetic activated chip-based separation (CMACS) device⁴⁰ and a micromagnetic separation (MMS) chip⁴¹ have also been reported. Both devices can precisely control the path of aptamer bound magnetic beads with high separation efficiency. However, there are limitations to each of these strategies. There are concerns regarding protein stability and integrity through the multiple selection cycles in the sol-gel-microheater device; and complicated target immobilization procedures, elongated incubation, and the potential for non-specific interactions with surface structures are of concern in the CMACS and MMS devices.

Capillary electrophoresis based selections (CE-SELEX) address many common issues encountered in traditional selections.⁴²⁻⁴⁵ Selections are performed in free solution, reducing the opportunity for non-specific interactions and eliminating complicated immobilization strategies. The high resolving power of CE increases the rate of enrichment allowing high affinity aptamers to be obtained in 2-4 rounds of selection. Unfortunately, CE-SELEX is not without limitations of its own. Only several nL of library can be injected without causing unacceptable loss of resolution.⁴⁵ This small volume limits the number of sequences that can be assessed and requires very high library concentrations. Fraction collection is also challenging since the abundance of aptamers in early rounds is often below the limit of detection and variability in mobilities requires collection windows to be adjusted on the fly.⁴⁶

Micro Free Flow Electrophoresis (μ FFE) provides a unique solution to the challenges presented by CE-SELEX. Contrary to many separations, μ FFE can be used to continuously introduce, separate and collect analytes (see Fig. 1).⁴⁷⁻⁵⁰ Analyte is continuously streamed into a planar separation chamber. An electric field is applied perpendicularly to the pressure driven flow, deflecting analytes laterally according to their mobility. μ FFE designs and separation conditions have been optimized allowing long term operation of the device.^{51, 52} μ FFE separation theory and sources of band broadening have been characterized.⁵³ Unique μ FFE detection strategies have been demonstrated.⁵⁴ Most interestingly, a gradient technique has been developed to measure aptamer-target equilibria, demonstrating the feasibility of using μ FFE to separate binding sequences from non-binding sequences.⁵⁵

In the current manuscript we report the first successful application of μ FFE in a SELEX selection for DNA aptamers. IgE was chosen as a target since aptamers have previously been selected for this protein using both traditional SELEX⁵⁶ and CE-SELEX^{42, 43} allowing a direct comparison with these approaches.

Experimental

Materials and Chemicals

Human myeloma IgE protein was purchased from Athens Research and Technologies (Athens, GA). Nuclease free H₂O, forward primer 5'-FAM-AGC AGC ACA GAG GTC AGA TG-3', reverse primer 5'-Biotin-TTC ACG GTA GCA CGC ATA GG-3', the initial ssDNA library 5'-FAM-AGC AGC ACA GAG GTC AGA TG(N)₄₀ CCT ATG CGT GCT ACC GTG AA-3', and the selected aptamers were from Integrated DNA Technologies, Inc. (Coralville, IA). For PCR reactions, dNTPs and the 25 bp DNA ladder were from Invitrogen (Carlsbad, CA); Taq polymerase and ThermoPol buffer were from New England BioLabs (Ipswich, MA); and the gel loading dye was from Promega (Madison, WI). Other chemicals

were purchased from Sigma Aldrich (St. Louis, MO) at the highest grade available, except Acetic acid (CH_3COOH , 99.7%, Mallinckrodt Baker), 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES, 99%, Alfa Aesar), MgCl_2 (99.8%, Mallinckrodt Baker), NaCl (99.0%, Spectrum), and KH_2PO_4 (99.9%, J. T. Baker). All buffers were prepared in nuclease free H_2O , and filtered through 0.2 μm membrane filters before use.

μFFE Fabrication

μFFE devices were fabricated using previously reported procedures.⁵⁵ Briefly, three photolithography steps were performed to define the electrode channels, the separation channel, and pattern the electrodes onto 1.1 mm borofloat wafer (Precision Glass & Optics, Santa Ana, CA) according to the design shown in Figure 1. The patterned chip was anodically bonded to a second borofloat wafer that had previously been drilled with inlet and outlet holes. Channel depths were approximately 20 μm in the separation channel and 100 μm in the electrode channels. The sample inlet hole had a diameter of 355 μm .

μFFE Separation

Before separation, the μFFE chip was coated with PEO to suppress the electroosmotic flow.⁵⁷ Briefly, 1 M HCl was first pumped into the chip at 6 mL/min for 10 min, followed by 0.2 % PEO in 0.1 M HCl at 3 mL/min for 10 min. Finally, the separation buffer (25 mM HEPES, 300 μM Triton X-100, adjusted to pH 7.0 by 1 M NaOH) was pumped at 6 mL/min for 10 min to remove HCl and unbound PEO. Before the selection, the ssDNA library was heated to 72 $^\circ\text{C}$ for 5 min and then gradually cooled to room temperature. For the 1st round of selection, 100 μM random sequence ssDNA library was incubated with 10 nM IgE at room temperature for 20 min in the binding buffer (tris(hydroxyamino)methane-glycine-potassium TKG) buffer, composed of 25 mM tris(hydroxyamino)methane, 192 mM glycien, and 5 mM KH_2PO_4 at pH 8.3). The following IgE concentrations were used in subsequent selection rounds: 10 nM in the 2nd round, 1 nM in the 3rd round, and 100 pM in the 4th round. The mixture (40 μL in total) was then loaded into a 100 μL syringe (Hamilton Company, Reno, NV), and pumped into the μFFE chip (Figure 1, hole (2)) using a syringe pump (PicoPlus, Harvard apparatus, Holliston, MA) at 100 nL/min. The separation buffer was pumped into the device (Figure 1, hole 1) using a second syringe pump (pump 22, Harvard Apparatus) at 1 mL/min. An electric field of 150 V/cm was applied across the chip so that the nonbinding ssDNA was separated from aptamer-IgE complexes. A stable current of 0.32 mA was observed over the 30 minute separation time. With EOF suppressed, the free ssDNA and the complex were split into two fractions (Figure 1, holes 3) and collected separately.

Laser Induced Fluorescence (LIF) Detection

The beam from a solid state laser (488 nm, 50 mW, Newport Corp, Irvine, CA) was expanded into a line and focused across the separation channel 1.5 cm downstream from the sample inlet. A microscope objective (3 \times zoom) was positioned above the detection zone. Fluorescence images were recorded every second using a Cascade 512B CCD camera (Photometrics, Tucson, AZ) through an AZ100 stereomicroscope (Nikon Corp., Tokyo, Japan). An Endow GFP bandpass emission filter cube (Nikon Corp., Tokyo, Japan) consisting of a dichroic mirror (495 nm cutoff) and two bandpass filters (450–490 nm and 500–550 nm) were used for wavelength selection. Images were processed using MetaVue software (Downington, PA). Linescans were also recorded using by MetaVue, and later analyzed in Cutter 7.0.⁵⁸

PCR Amplification and Purification

Collected sequences were PCR amplified immediately after each selection round. In the final reaction vials, there were 1 mM each of the four dNTPs, 7.5 mM MgCl₂, 500 nM forward and reverse primers, 1 × ThermoPol buffer and 0.05 units/μL *Taq* polymerase. A negative control with all PCR reagents but no ssDNA was also performed every round to verify the absence of background contamination. 23 cycles of denaturation (94 °C, 30 s), annealing (55 °C, 30 s), and extension (72 °C, 20 s) were performed, with a final extension at 72 °C for 5 min. A 1.5 % agarose gel with ethidium bromide staining was used to confirm the presence of the desired PCR products. Single stranded FAM labeled DNA sequences were obtained using an on column purification followed by the ethanol precipitation.⁴³

Dissociation constant K_d measurements

Affinities of the selected pools and individual aptamer sequences were measured using both affinity capillary electrophoresis (ACE) and fluorescence polarization (FP). Approximately 2.5 nM ssDNA samples were titrated with increasing concentrations of human IgE. A commercial CE system (P/ACE MDQ, Beckman Coulter, Inc., Fullerton, CA) equipped with laser induced fluorescence detection ($\lambda_{\text{ex}} = 488 \text{ nm}$, $\lambda_{\text{em}} = 520 \text{ nm}$) was used to perform ACE experiments. Samples were injected into a 50 cm fused silica capillary (Polymicro Technologies, Phoenix, AZ) at 1psi for 4 sec. The separations were then performed at 30 kV in TGK buffer. The peak heights of the unbound ssDNA were used to calculate the bound fractions and estimate K_d using the following equation:⁵⁹

$$f_a = \frac{c}{1 + K_d / ([P]_t - 0.5([D]_t + [P]_t + K_d - (([D]_t + [P]_t + K_d)^2 - 4[D]_t[P]_t)^{0.5}))} \quad (2)$$

in which f_a is the bound fraction, $[P]_t$, $[D]_t$, and c are total IgE concentration, total DNA concentration, and maximum bound fraction, respectively. $[P]_t - 0.5([D]_t + [P]_t + K_d - (([D]_t + [P]_t + K_d)^2 - 4[D]_t[P]_t)^{0.5})$ represents the free IgE concentration.

In FP experiments, 15 μL of the same samples used in ACE were loaded into a corning 3540 microplate (Corning Incorporated, Corning, NY) and experiments were performed on a Synergy™ 2 Microplate Reader (BioTek Instruments, Inc., Winooski, VT). Parallel and perpendicular intensities ($\lambda_{\text{ex}} = 485 \pm 20 \text{ nm}$, $\lambda_{\text{em}} = 528 \pm 20 \text{ nm}$) were measured, and polarization values were calculated using Gen 5™ software (BioTek Instruments, Inc., Winooski, VT). The bound fraction was determined according to:

$$f_a = \frac{P - P_o}{P_m - P_o} \quad (3)$$

in which P , P_o and P_m are the measured polarizations of a sample, free DNA, and the complex, respectively. Meanwhile, the overall fluorescence intensity was monitored and corrections to the bound fractions were made according to the intensity change as previously reported.⁶⁰

DNA cloning and sequencing

A TOPO TA Cloning® Kit for Sequencing (Invitrogen, Carlsbad, CA) was used to perform cloning reactions and grow colonies. Briefly, 1 μL of the selected ssDNA from pool 1 through pool 4 was added to 99 μL PCR master mixes and amplified separately, as described above, except using unlabeled forward and reverse primers. 4 μL of the 100 μL fresh PCR products were incubated with TOPO® vector at room temperature for 5 min to allow the cloning reactions to take place. The inserted vectors were then chemically transformed into

One Shot[®] TOP10 competent *E.coli* cells. The mixtures were incubated in a 37 °C shaking incubator for 30 min and then spread onto selective plates containing 50 µg/mL kanamycin. For each pool, two agar plates were used to grow colonies, which produced 300 to 400 colonies. Individual colonies were then randomly picked and cultivated in a 96 well plate with liquid LB containing 50 µg/mL kanamycin at 37 °C for another day. Cell pellets were sent to the Biomedical Genomics Center at the University of Minnesota for Sanger sequencing. Since some wells of the plate were used as controls, and some pellets were not sufficient to obtain confident sequencing information, 23, 28, 19, and 6 sequences were obtained from pools 1, 2, 3, and 4, respectively (See Table S1 for aptamer sequences).

Results and discussion

µFFE selection

Figure 2 demonstrates the µFFE separation observed during the first round of selection. An electric field of 150 V/cm was applied across the flow channel. Unbound ssDNA sequences were deflected toward the anode due to the suppressed electroosmotic flow (EOF). The aptamer-IgE complexes, which were only deflected minimally, were well separated from the unbound sequences. It should be noted that µFFE facilitates collection of analytes with low mobilities. Pressure rinses were necessary to observe²⁴ or collect^{42, 43} aptamer-IgE complexes in previously reported CE separations. The flow was split into two streams at the exit of the µFFE separation chamber. The IgE-DNA complexes were collected into a centrifuge tube through the tubing connected to the right nanopore. This continuous collection strategy was much simpler than that of CE-SELEX, eliminating the complicated timing associated with collecting complexes as they migrated off the end of a capillary. A linescan (Figure 2b) recorded across the detection zone reinforces how well resolved the binding sequences are from the non-binding sequences. The high concentration of library (100 µM) used in the first round of selection caused the signal for the unbound sequences to go off scale but this was not a concern since the primary goal was to identify the position of the peaks for accurate fraction collection, not quantitation.

In the first round of selection, 100 µM of fluorescently tagged ssDNA library was incubated with 10 nM IgE in TGK binding buffer for 20 min. After incubation, 3 µL of the library-IgE mixture was introduced into the µFFE device over a period of 30 minutes, which corresponded to 1.8×10^{14} sequences. A typical CE separation only allows a discrete injection of approximately 8 nL.⁴⁵ Over 300-fold more volume, and therefore sequences, could be introduced into the µFFE device. The number of sequences could easily be increased further by increasing the analyte flow rate, library concentration or collection time.

The multiple-depth µFFE design⁵¹ generated high flow over the electrodes to dissipate electrolysis bubbles while minimizing flow, and therefore dilution, in the separation chamber. The dilution factor is defined as the ratio of the collected sample volume to the introduced sample volume. In a µFFE device, the planar flow profile can be described using lubrication theory according to the following equation:⁶¹

$$q = \Delta P H^3 w / 12 \eta L \quad (1)$$

in which q is the volumetric flow rate, ΔP is the pressure difference, H is the channel depth, w is the channel width, η is the buffer viscosity, and L is the channel length. Comparing the electrode and separation channels: the ratio of channel depths is $20 \mu\text{m} / 100 \mu\text{m} = 0.2$ (separation channel/electrode channel), the ratio of channel widths is $10 \text{mm} / (2 \times 2 \text{mm}) = 2.5$ (i.e. two electrode channels), and the ratio of channel lengths is approximately 1. The ratio of flow volume through the channels is $0.2^3 \times 2.5 / 1 = 1:50$ (separation channel/

electrode channel). Solution is only collected from half of the separation channel. As result, approximately 1% of the total volume through the μ FFE device is collected. Over a period of 30 min, 30 mL separation buffer and 3 μ L sample are introduced into the chip, resulting in a collected volume of 300 μ L. This calculation agrees very well with experimental results. Comparing with the 3 μ L of library injected yields a modest 100-fold dilution of the binding sequences during the μ FFE separation. This compares favourably with dilution factors of 6,000 typically encountered in CE-SELEX.^{42–45} The higher DNA concentration in the collected fraction and increased volume collected facilitates PCR and decreases the potential for contamination or non-specific amplification. Furthermore, the transit time through the μ FFE flow chamber is only 10–20 seconds, greatly decreasing the potential for dissociation during the separation when compared to CE, in which separations typically take 5–15 minutes. The electric field applied in μ FFE (150 V/cm) is also lower than that typically used in CE-SELEX (~500 V/cm), further decreasing the potential for dissociation.

Aptamer Characterization

Four rounds of μ FFE selection were performed. The affinity of the selected pools for IgE was monitored using two orthogonal methods⁶²: fluorescence polarization (FP)⁶⁰ and affinity capillary electrophoresis (ACE)⁴² (see Experimental, Dissociation constant K_d measurements). As shown in Figure 3, the initial library had a low affinity for IgE in TGK buffer, ($K_d = 460 \pm 160$ nM and 550 ± 280 nM as measured by FP and ACE, respectively). A significant improvement in affinity was observed after a single round of μ FFE selection, with K_d values of 29 ± 15 nM and 58 ± 55 measured by FP and ACE, respectively. No further improvement in affinity was observed after the second round of selection. IgE concentration was decreased by a factor of 10 (1 nM) in the third round of selection and by a factor of 100 (100 pM) in the fourth round of selection in an effort to improve the stringency of the selection. However, the enhanced stringency did not result in improved affinity for IgE, suggesting that the selection had converged after a single round of μ FFE selection.

Aptamers were randomly cloned and sequenced from the DNA pools after each round of selection. 23 sequences were identified from round one, 28 sequences from round two, 19 sequences from round three, and 6 sequences from round four (see supplementary information for aptamer sequences). These sequences were analyzed to identify homologous sequences or motifs. As shown in Table 1, no identical or similar sequences were found in round 1; two identical sequences were found in rounds 2 and 3; three identical sequences were found in round 4 with a fourth sequence demonstrating 89% similarity as revealed by software ClustalW2[®]. It should be noted that sequences that appeared multiple times in one round did not share any conserved region or motifs when compared with sequences isolated from other rounds. Although the number of the clones was limited, identification of identical sequences in such a small sample size suggests a decrease in diversity as the pool converged. This is particularly evident in round four, where 4 out of 6 sequences isolated were very similar. The affinities of all sequences identified multiple times were measured by FP and ACE (see Table 1). The measured dissociation constants were remarkably similar and agreed well with the affinities measured for the bulk pools.

Finally, three aptamers were chosen from the cloned sequences obtained from rounds one and three using a random number generator. Dissociation constants for each of these sequences are shown in Table 1 (see Figure 4 for representative binding curves).

No statistical difference was observed between sequences obtained after one or three rounds suggesting that the selection had converged after a single round of μ FFE selection. K_d values for sequences chosen randomly were statistically indistinguishable from those that appeared multiple times in the cloning results. This result is similar to previous CE-SELEX selections that yielded seemingly diverse pools of aptamers with similar affinities.^{42, 43} It

should be noted that every sequence assessed demonstrated low nM affinity for IgE, even in the absence of negative selections, suggesting that similar to CE-SELEX, performing selections in free solution diminishes the opportunity for non-specific interactions. Dissociation constants measured using ACE were consistently higher than those measured using FP. Yang *et al* have demonstrated that the high electric field of CE can affect observed half-lives and binding constants of protein complexes.⁶³ This raises a concern regarding whether the high field typically used in CE-SELEX and whether this field modifies the selection environment, affecting the applicability of the selected aptamers. Buchanan *et al*⁶⁴ have demonstrated that a combination of low electric field and shorter time in the field minimize dissociation of aptamer-target complexes during a CE separation. In practice this can be difficult to balance in CE since separation time is inversely proportional to electric field. μ FFE does not share this difficulty since electric field and separation time can be optimized independently. This unique property of μ FFE is advantageous allowing more control over separation conditions when compared to CE.

Aptamers for IgE obtained using μ FFE demonstrate similar dissociation constants to those obtained previously using CE-SELEX (~20 nM)^{42,43} and conventional SELEX (~10 nM)⁵⁶. It should be noted that these aptamers were obtained after a single round of μ FFE selection, which compared favorably with the 2–4 rounds required by CE-SELEX^{42, 43} or the 15 rounds required by conventional SELEX.⁵⁶

Conclusions

In conclusion, we demonstrated the advantages of isolating aptamers using μ FFE selections. Within 30 min of continuous μ FFE separation and collection, 1.8×10^{14} sequences were assessed, a 300-fold improvement over CE-SELEX. μ FFE also eliminated the complicated timing associated with fraction collection in CE-SELEX. Four selection cycles were completed within four days. Low nanomolar affinity sequences were identified after a single round of μ FFE selection, suggesting that aptamers could be obtained even faster. Although the device was expected to have lower separation efficiency than CE, obtaining high affinity aptamers after a single selection round suggests a high rate of enrichment was achieved. The free solution μ FFE separation simplifies the protocol, eliminating the need for target immobilization, elongated incubation, or negative selections. With these advantages, we believe that μ FFE device can be readily adopted to select aptamers for a wide range of targets.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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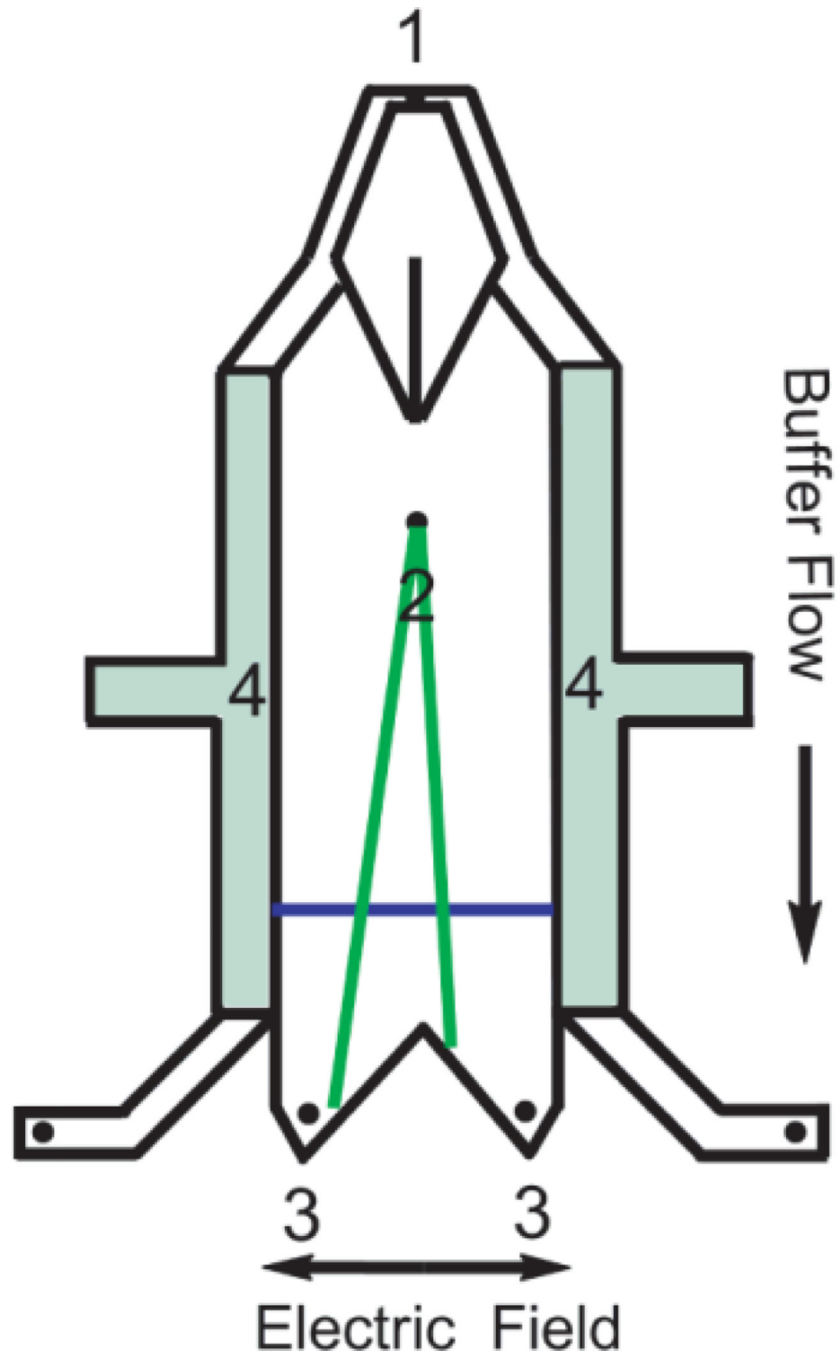


Figure 1. Schematic of a μ FFE device demonstrating the buffer inlet (1), sample inlet (2), fraction collection outlets (3) and electrode channels (4). The blue line denotes the detection zone where the laser line is expanded across the separation channel.

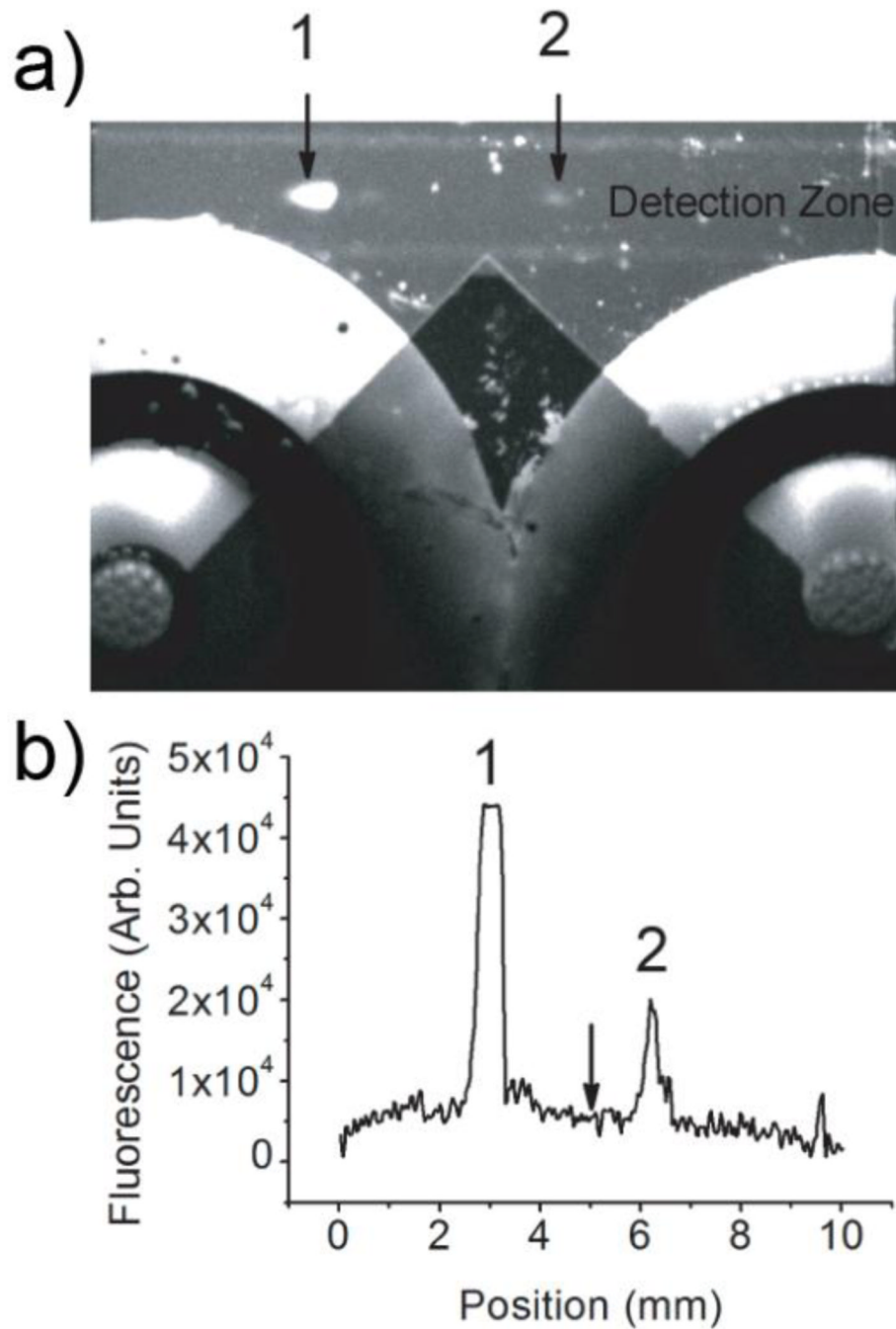


Figure 2.

(a) An image of a μ FFE separation of free (1) and bound ssDNA (2) recorded during selection cycle #1. The fraction collection channels and outlet ports are clearly visible in this image. (b) A linescan across the detection zone imaged in (a). The arrow indicates the fraction cutoff point at the exit of the μ FFE channel demonstrating clear separation of free (1) from bound (2) ssDNA. The anode is on the left in all images.

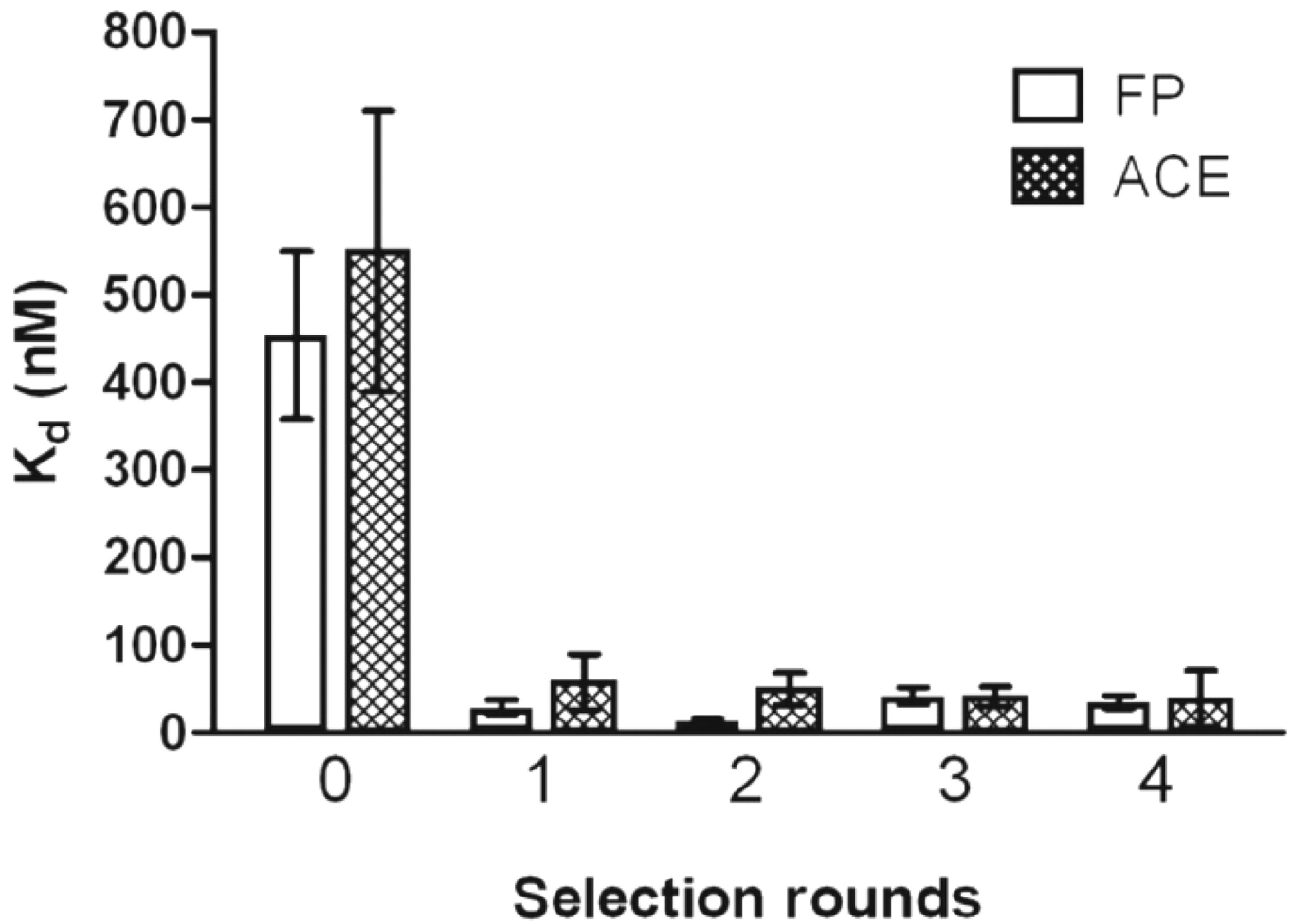


Figure 3. K_d of the starting library and the selected pools measured by fluorescence polarization (FP) and affinity capillary electrophoresis (ACE). Error bars represent 95% confidence interval.

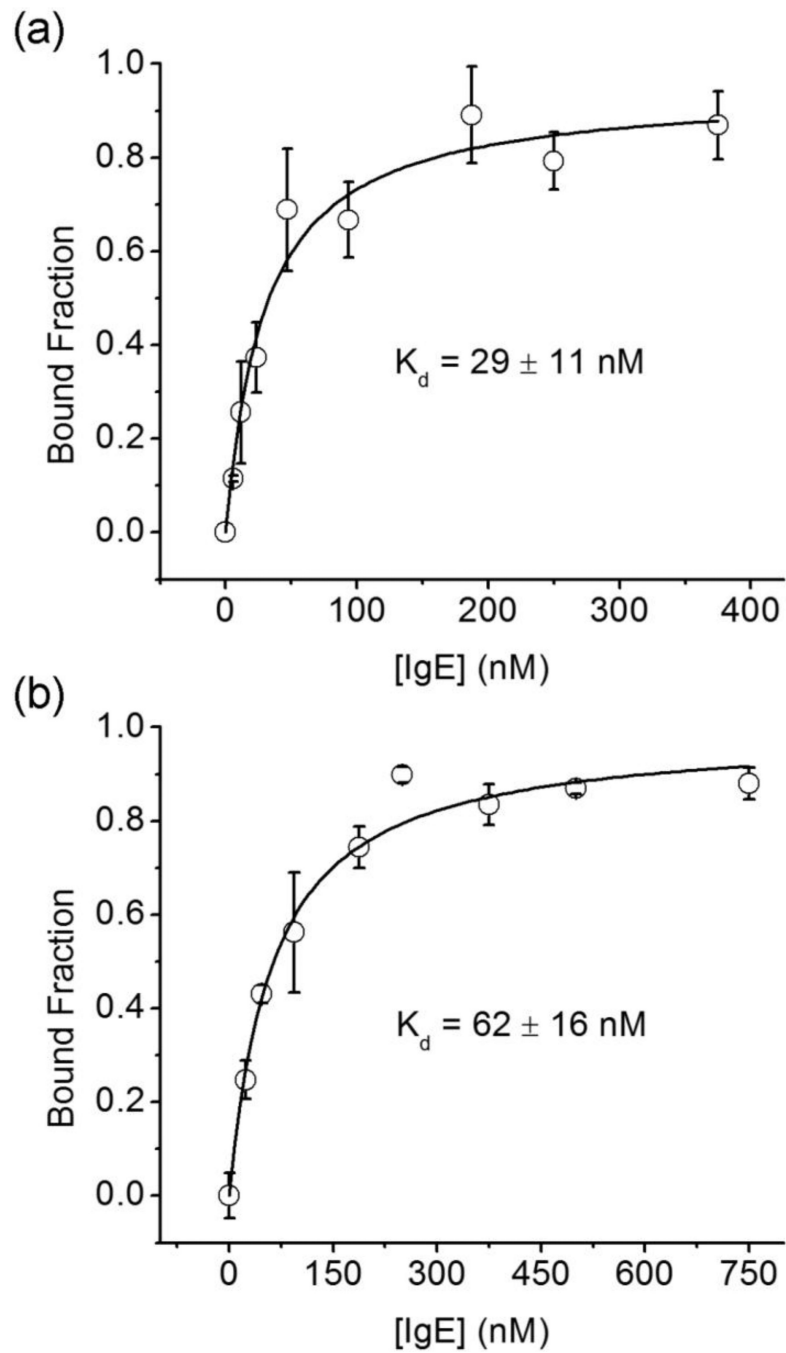


Figure 4. Binding curves for sequence 3.9, obtained using (a) FP and (b) ACE. Three measurements were taken at every IgE concentration. Error bars represent standard deviation. Errors of the K_d values represent 95% confidence interval.

Table 1 K_d of selected aptamers

Aptamer ^{a)}	FP K_d ^{b)}	ACE K_d ^{b)}
<i>Multiple copy sequences identified</i>		
Clone 2.13 and 2.24	20 ± 4 nM	136 ± 58 nM
Clone 3.5 and 3.14	29 ± 7 nM	33 ± 21 nM
Clone 4.4, 4.5, and 4.6	39 ± 20 nM	58 ± 17 nM
Clone 4.2	44 ± 15 nM	62 ± 29 nM
<i>Randomly chosen sequences from round 1 and round 3</i>		
Clone 1.4	23 ± 6 nM	58 ± 33 nM
Clone 1.13	22 ± 6 nM	63 ± 17 nM
Clone 1.18	32 ± 11 nM	66 ± 35 nM
Clone 3.2	20 ± 7 nM	50 ± 27 nM
Clone 3.9	29 ± 11 nM	62 ± 16 nM
Clone 3.11	28 ± 13 nM	47 ± 17 nM

^{a)} Aptamers were given unique identifiers where the first number gives the selection round and the second number identifies a particular sequence cloned in that round.

^{b)} Errors represent 95% confidence interval.