Characterization of linearly coupled capillaries with various inner diameters in the context of capillary electrophoresis

Daniel Böhm1 · Frank-Michael Matysik1

Received: 28 January 2021 / Accepted: 5 May 2021 © The Author(s) 2021

Abstract
As a result of continuous instrumental progress, capillary electrophoresis has become an established separation technique. However, the choice of the suitable capillary inner diameter is sometimes difficult due to different instrumental requirements concerning injection, separation, or detection. To overcome this problem, we assembled two capillaries with different inner diameters, meaning that the inner diameter of the capillary at the injection side was different from that at the detection side. Since this was a rather uncommon approach, we focused on the associated effects in this proof-of-concept study. For the experiments, a non-aqueous model system was used, consisting of an acetonitrile-based background electrolyte and the two ferrocene derivates, ferrocenemethanol and decamethylferrocene. Using capillary flow injection analysis hyphenated to capacitively coupled contactless conductivity detection, it could be shown that fragmented capillaries of the same inner diameter had slightly lower volume flow rates than non-fragmented capillaries. With non-aqueous capillary electrophoresis hyphenated to UV detection, it was found that the coupling of capillaries with different inner diameter had a much stronger effect on the capillary electrophoresis flow than combinations with the same inner diameter. Additionally, if the inner diameter of the second capillary was larger than the inner diameter of the first capillary, a higher theoretical plate number and an increased sensitivity were found. Furthermore, it was found that there was no significant peak tailing introduced by the coupling.

Graphic abstract

Keywords Capillary flow injection analysis · Capillary zone electrophoresis · Coupled fused silica capillaries · Electrochemistry · Organometallic compounds

Introduction
Capillary electrophoresis (CE) is a powerful separation technique due to its high separation efficiency, the low sample consumption, and its high separation speed. Thus, the field of CE is still in instrumental progress [1–3]. In recent years, different injection methods were developed and CE was hyphenated to various detectors [4, 5]. However, choosing the suitable capillary inner diameter (ID) is sometimes difficult. Capillaries with small IDs are for example interesting for small amounts of sample and high separation efficiency due to lower Joule heating [4, 6, 7]. In contrast, capillaries with larger IDs are preferred for example for on-column UV/Vis detection to generate a better sensitivity due to extended optical path lengths [4, 8]. These are just a few examples which show that different applications require...
various capillary IDs. When injection, separation or detection aspects suggest the use of different capillary ID, often a compromise has to be found. We tried to overcome this problem by the combination of two capillaries with various IDs.

The coupling of capillaries is already described in literature but predominately in the context of 2D separations. In these cases, the capillaries were coupled with valves or micro-structured chips [9–13]. These approaches were not suitable for our application because we intended to have a continuous CE separation without decoupling from the high voltage field and a straightforward combination of conventional capillaries. There are other cases where capillaries with different IDs were coupled to achieve fast electrophoretic separations [14–16]. However, in these studies, injection and detection were performed in capillary segments of identical ID. Horká et al. [17] used capillaries with different IDs produced by etching for online sample pre-concentration. In most of the previously mentioned cases, the capillaries were irreversibly connected. In our approach, a commercially available capillary connector was used, which allowed for exchanging of capillaries and nearly dead volume-free coupling. Additionally, injection and detection were performed on opposite ends of the linearly coupled capillaries.

In the frame of this contribution, we investigated the linear coupling of capillaries with various IDs, meaning that the ID of the capillary at the injection side was different from that at the detection side. The resulting effects were investigated with capillary flow injection analysis hyphenated to capacitively coupled contactless conductivity detection (CFIA-C4D) and CE hyphenated to UV detection (CE-UV).

Results and discussion

CFIA-C4D experiments

The measurements were performed with a model system containing decamethylferrocene (dMFc) as analyte and a non-aqueous, acetonitrile-based background electrolyte (BGE) as carrier solution. This model system was chosen due to the good UV transparency and the stable electroosmotic flow. Additionally, in future projects, it is planned to utilize the concept of coupled capillaries in combination with an amperometric detector, where the acetonitrile-based BGE is favorable due to the electrochemical characteristics [4, 18].

The CFIA-C4D experiments in this project were performed to study the effects associated with capillary coupling under conditions of a gravity-driven flow system. The volume flow rates (VFRs) and the corresponding standard deviations (SDs) shown in Table 1 for a height difference of 20 cm were calculated based on four CFIA-C4D measurements. It should be noted that different linear flow velocities were present in different segments of the fragmented capillaries with various IDs. Therefore, we used the VFRs which were the same over the entire length of the capillary. It was found that the VFRs were slightly lower for fragmented capillaries compared to non-fragmented capillaries of the same dimension. This indicates that a small additional flow resistance arises when two capillaries are combined. Furthermore, it was found that the VFR decreases for upstream capillaries with lower IDs and vice versa. The VFR for the combination 75 + 25 µm could not be determined due to bubble formation.

CE-UV experiments

Figure 1 shows in the upper part the electropherograms of the non-fragmented capillaries with IDs of 25, 50, and 75 µm and in the lower part the electropherograms of the capillary combinations with a downstream 50 µm capillary segment (25 + 50, 50 + 50, and 75 + 50 µm). Prior to the detailed discussion, it has to be mentioned the neutral ferrocenemethanol (FcMeOH) served as electroosmotic flow (EOF) marker while dMFc served as cationic model

| Capillary combination/µm | VFR/10⁻⁹ dm³/s | ± SD/10⁻⁹ dm³/s |
|--------------------------|----------------|----------------|
| 25                       | 0.0577         | 0.0005         |
| 25 + 25                  | 0.057          | 0.001          |
| 50 + 25                  | 0.0775         | 0.0007         |
| 75 + 25                  | -a             | -a             |
| 25 + 50                  | 0.137          | 0.005          |
| 50                       | 0.865          | 0.002          |
| 50 + 50                  | 0.80           | 0.05           |
| 75 + 50                  | 1.05           | 0.06           |
| 25 + 75                  | 0.187          | 0.002          |
| 50 + 75                  | 1.941          | 0.003          |
| 75                       | 4.56           | 0.02           |
| 75 + 75                  | 4.2            | 0.1            |

Experimental conditions: sample solution 10 mmol/dm³ dMFc in BGE, capillary length 70 cm (40 cm to the detector, fragmented capillaries: 20 cm first part and 50 cm second part), injection time 5 s and BGE (10 mmol/dm³ NH₄OAc and 1 mol/dm³ HOAc in ACN) as carrier solution

*a For the capillary combination 75 + 25 µm, no VFR could be determined due to bubble formation
Characterization of linearly coupled capillaries with various inner diameters in the context…

analyte since it is easily oxidized by dissolved oxygen in solution [19]. The BGE levels in the inlet and outlet reservoirs were at the same height. Therefore, there was no gravity-driven flow. The measurements performed with non-fragmented capillaries had nearly the same migration times for the cationic (dMFc⁺) and the neutral species (FcMeOH), independent of the ID. The averaged migration times and the corresponding SDs for the non-fragmented and fragmented capillaries are listed in Table 2. Since the electrophoretic and the EOF migration contribute to the overall migration time of the dMFc⁺, we calculated the apparent mobility of dMFc⁺ (µa), the EOF mobility of FcMeOH (µEOF), and the effective mobility of dMFc⁺ (µeff) to separate these effects. Due to the very similar migration times for the non-fragmented capillaries (25, 50, and 75 µm), the mobilities were nearly the same. The relative standard deviations for the mobilities of the non-fragmented capillaries with different IDs were 3% for µa, 6% for µEOF and 2% for µeff (n = 9). The calculated mobilities (µa, µEOF, and µeff) and the corresponding SDs for the different capillary combinations are summarized in Table 3.

The electropherograms for the capillary combinations with a downstream 50 µm capillary segment (25 + 50, 50 + 50, and 75 + 50 µm) are presented in the lower part of Fig. 1. For the combination 50 + 50 µm, the migration times were 4% higher for the cationic (dMFc⁺) and 7% higher for the neutral species (FcMeOH) compared to the non-fragmented 50 µm capillary. Looking at the mobilities, it was found that µEOF was 8% and µeff was 1% lower for the fragmented capillary (50 + 50 µm). Therefore, the coupling of two identical capillaries predominately affected the EOF. The reduction of the EOF was probably due to a flow resistance at the connection site, which was also found for the CFIA-C⁴D experiments.

In contrast to the combination of capillaries with the same ID, a considerable shift of the migration times and the corresponding mobilities was found for the fragmented capillaries with different IDs (25 + 50 and 75 + 50 µm). Since µEOF and µeff were significantly lower for fragmented capillaries with different ID, it can be concluded that the coupling influenced the EOF and the migration of the cationic species. It is important to note that in the present case, µeff does

---

Table 2: CE-UV studies for the determination of the migration times for dMFc⁺ and FcMeOH, the measured electrophoretic currents and the corresponding SDs (n = 3) of non-fragmented and fragmented capillaries

| Capillary combination/µm | Migration time dMFc⁺/s ± SD/s | Migration time FcMeOH/s ± SD/s | Electrophoretic current/µA ± SD/µA |
|--------------------------|-------------------------------|-------------------------------|-----------------------------------|
| 25                       | 128 ± 1                       | 319 ± 2                       | 1.42 ± 0.01                       |
| 50                       | 120.1 ± 0.9                   | 281 ± 3                       | 3.98 ± 0.09                       |
| 75                       | 120.7 ± 0.2                   | 295 ± 1                       | 9.7 ± 0.1                         |
| 25 + 50                  | 146.3 ± 0.3                   | 379 ± 1                       | 2.92 ± 0.08                       |
| 50 + 50                  | 124 ± 2                       | 300 ± 3                       | 4.02 ± 0.01                       |
| 75 + 50                  | 180 ± 1                       | 439 ± 4                       | 4.84 ± 0.08                       |

Experimental conditions: 1 mmol/dm³ FcMeOH and dMFc in BGE (10 mmol/dm³ NH₄OAc and 1 mmol/dm³ HOAc in ACN), injection segment 0.35 cm, separation voltage 25 kV, capillary length 70 cm (40 cm to the detector, fragmented capillaries: 20 cm first part and 50 cm second part), UV detection at 210 nm.
the combination 75 + 50 µm by the narrower downstream capillary. This explains why the species for the combination 75 + 50 µm showed lower mobilities in comparison to the capillary combination 25 + 50 µm.

To prove this hypothesis, CE-UV experiments with the inverted capillaries (upstream 50 µm capillary and downstream 25, 50, or 75 µm capillary) were made. According to the hypothesis, the capillary combination 50 + 25 µm should have the lowest mobilities, because in this configuration, the CE flow hits the front surface of the 25 µm capillary. The CE-UV experiments with the reverse capillary combinations confirmed this assumption because the lowest mobilities were found for the capillary combination 50 + 25 µm (electropherograms not shown). Based on these experiments, it can be concluded that the coupling influenced the CE flow due to a mechanical hindrance at the connection site. Especially for capillary combinations starting from a larger capillary ID and going to a smaller ID, a decrease of the mobilities was found.

Another striking point was that the peak areas for the combination 25 + 50 µm were about 2 times higher than the peak areas for the non-fragmented 25 µm capillary, although the same injection volume was applied in both cases (electropherograms shown in Fig. 1). This can be explained by Lambert–Beer law because with a downstream 50 µm capillary the layer thickness is two times higher than with a 25 µm capillary. Additionally, the lower linear flow velocity in the downstream 50 µm capillary compared to the upstream 25 µm capillary contributed to the increased peak areas. Due to the slower linear flow velocity, the analytes spent more time in the detection window, which led to higher peak areas.

Furthermore, it was demonstrated that all peaks showed nearly Gaussian shape and for all combinations there was only negligible peak tailing. Additionally, there was an increase of the theoretical plate number found for the capillary combinations starting from a smaller ID and going to a

---

**Table 3** Calculated mobilities $\mu_a$ for dMFc$^+$, $\mu_{EOF}$ for FcMeOH, $\mu_{eff}$ for dMFc$^+$, and the corresponding SDs ($n = 3$) of non-fragmented and fragmented capillaries based on the CE-UV experiments

| Capillary combination/µm | $\mu_a$ dMFc$^+$/cm$^2$ kV$^{-1}$ s$^{-1}$ | $\pm SD$/cm$^2$ kV$^{-1}$ s$^{-1}$ | $\mu_{EOF}$ FcMeOH/cm$^2$ kV$^{-1}$ s$^{-1}$ | $\pm SD$/cm$^2$ kV$^{-1}$ s$^{-1}$ | $\mu_{eff}$ dMFc$^+$/cm$^2$ kV$^{-1}$ s$^{-1}$ | $\pm SD$/cm$^2$ kV$^{-1}$ s$^{-1}$ |
|--------------------------|---------------------------------|-----------------|-----------------|-----------------|---------------------------------|-----------------|
| 25                       | 0.873 0.008 0.351               | 0.002 0.522     | 0.006           |
| 50                       | 0.933 0.007 0.398               | 0.005 0.535     | 0.002           |
| 75                       | 0.928 0.001 0.380               | 0.002 0.5484    | 0.0006          |
| 25 + 50                  | 0.765 0.002 0.295               | 0.001 0.4701    | 0.0008          |
| 50 + 50                  | 0.90 0.01 0.374                | 0.004 0.527     | 0.007           |
| 75 + 50                  | 0.623 0.004 0.255               | 0.003 0.368     | 0.001           |

$^a$$\mu_a$ was calculated according to the equation $\mu_a = (l/L)(t_m/V)$, where $l$ was the effective capillary length to the detector, $L$ was the total capillary length, $t_m$ the migration time of the analyte ion, and $V$ the applied voltage [20].

$^b$$\mu_{EOF}$ was calculated according to the equation $\mu_{EOF} = (l/L)(t_{EOF}/V)$, where $t_{EOF}$ was the migration time of the EOF marker [20].

$^c$$\mu_{eff}$ was calculated according to the equation $\mu_{eff} = \mu_a - \mu_{EOF}$ [20].

---

Fig. 2 Scheme of the capillary combinations with a downstream 50 µm capillary at the connection site: 25 + 50 (a), 50 + 50 (b), and 75 + 50 µm (c). Especially for the capillary combination 75 + 50 µm, the CE flow was hindered by the downstream smaller capillary.
larger ID. The theoretical plate numbers for the dMFc$^+$ and FcMeOH peak were $27 \times 10^3 \pm 3 \times 10^3$ and $28 \times 10^3 \pm 5 \times 10^3$ for the capillary combination 25 + 50 µm while the theoretical plate numbers for the non-fragmented 50 µm capillary were $81 \times 10^2 \pm 3 \times 10^2$ for dMFc$^+$ and $206 \times 10^2 \pm 9 \times 10^2$ for FcMeOH. This resulted from a compression of the peaks at the transition between a capillary starting with a smaller ID and going to a capillary with a larger ID. This effect could be explained with the different linear flow velocities in the capillary combination 25 + 50 µm. The linear flow rate in the upstream 25 µm capillary was higher than the linear flow velocity in the downstream 50 µm capillary, which led to the compression of the peaks at the connection site.

We tried to visualize this by the behavior of a dye plug at the connection site between a capillary with a smaller and a larger ID. For this purpose, a transparent capillary connector was fabricated. Figure 3 shows a series of microscopic images from a dye plug at the connection site between a 25 and 75 µm capillary. Figure 3c shows the long dye segment in the 25 µm capillary which shortens at the transition to the 75 µm capillary. The microscopic image 3e shows the compressed dye plug in the 75 µm capillary piece. Even if this experiment was performed by a simple CFIA setup, it could explain the increase of the theoretical plate number by the compression of the peaks at the connection site.

With the findings listed in this manuscript, the questions pointed out in the introduction could be clarified. It could be shown that CE measurements with linearly coupled capillaries of various IDs could be realized. Additionally, an increased sensitivity was found for on-column UV detection for the capillary combination 25 + 50 µm, due to the extended optical path length. The same capillary combination (25 + 50 µm) showed also a higher theoretical plate number than a non-fragmented 50 µm capillary. Contrary to expectations, no significant changes in the peak shape or peak tailing were found for capillary combinations with the same or different IDs. From the CFIA and the CE measurements, it could be concluded that there was a mechanical disturbance of the flow due to the coupling. Unlike the capillary combination with the same ID, a shift towards lower mobilities was found for capillary combinations starting from a larger ID and going to a smaller ID which was a drawback of fragmented capillaries.

Based on the results described in this manuscript, it can be concluded that the concept of linearly coupled capillaries with various IDs could be established. Therefore, it is possible to combine two capillaries with different IDs to adapt capillary dimensions to the respective injection, separation, or detection requirements.

**Conclusion**

To overcome the problem of choosing the best-suited capillary ID for injection, separation, or detection in CE, the concept of fragmented capillaries with various IDs was introduced and a proof-of-concept study was carried out. With this approach, it is possible to select the ideal capillary ID combination for the respective application.

**Experimental**

The following chemicals were used, all of analytical grade: ferrocenemethanol and decamethylferrocene (purity 99%, ABCR, Karlsruhe, Germany), acetonitrile (ACN), ammonium acetate (NH$_4$OAc), 0.1 mol/dm$^3$ sodium hydroxide solution, ultrapure water provided by a MilliQ Advantage A10 system (Merck, Darmstadt Germany),

---

**Fig. 3** Subsequently recorded microscopic images (a–e) showing the behavior of a dye plug at the connection site between a 25 (left side) and a 75 µm (right side) capillary. The photos show the compression of the dye plug at the transition between a capillary with a smaller and a capillary with a larger ID. Experimental parameters: dye solution (methylene blue in BGE), the flow was gravity-driven by BGE (10 mmol/dm$^3$ NH$_4$OAc and 1 mmol/dm$^3$ HOAc in ACN).
acetic acid (HOAc; Roth, Karlsruhe, Germany), methylene blue (Riedel-de Haën, Seelze, Germany).

Capillaries

For both experiments (CFIA-C 4D and CE-UV), capillaries with IDs of 25, 50, and 75 µm, an outer diameter of 360 µm, and a total length of 70 cm were used. They were purchased from Polymicro Technologies (Phoenix, USA). Measurements were carried out with fragmented and non-fragmented capillaries. For the measurements with the fragmented capillaries, the original capillaries were cut into two pieces (20 and 50 cm) yielding a total of 9 possible capillary combinations with a total length of 70 cm (20 cm first capillary piece and 50 cm second capillary piece). These combinations are summarized in Table 1. At both ends of the capillaries, about 2 mm of the polyimide coating was removed to avoid a dead volume introduced by polyimide swelling. To enable a nearly dead volume-free coupling, both sides of the capillary pieces were polished to receive planar capillary tips. For the linear assembling of the capillaries, both capillary pieces were placed in Micro Tight Sleeves F 185 and linearly assembled in a capillary connector called Union Assembly Micro Tight P 720 both from IDEX Health & Science (Oak Harbor, USA). The installation of the capillaries was carried out according to the user manual of the capillary connector. A detailed scheme of the two capillaries assembled in the capillary connector is shown in Fig. 4. Before the first CE measurements, the capillaries were preconditioned by flushing them for 10 min with 0.1 mol/dm³ sodium hydroxide solution, afterwards for 5 min with ultrapure water, and in the end for 30 min with BGE (10 mmol/dm³ NH₄OAc, 1 mol/dm³ HOAc in ACN).

CFIA-C 4D setup

The VFRs for the fragmented and non-fragmented capillaries were determined with a CFIA-C 4D setup schematically depicted in Fig. 5. The concept of CFIA with gravity-driven flow was first described by Matysik et al. [21]. The flow in the capillary was gravity-driven by a permanent height difference \( \Delta h \) between the inlet and outlet reservoir. We always used the term BGE instead of carrier solution because both solutions were identical. The injection time for all CFIA-C 4D experiments was 5 s. A laboratory-constructed autosampler of a CE device was used for the hydrodynamic injection. The sample solution consisted of 10 mmol/dm³ dMFc in BGE. A high-resolution C 4D was placed after 40 cm from the injection side for detection. The detector described elsewhere was constructed in the group of Prof. C. L. do Lago (São Paulo, Brazil) [22]. A double determination at two different heights was done for the calculation of the VFRs.

CE-UV setup

Figure 5 shows a scheme of the CE-UV setup on the right side. It consisted of a laboratory-constructed CE device with an integrated autosampler. This device was connected to a T1CP 300 304p high voltage power supply from ISEG (Radeberg, Germany). The whole CE system was placed in an acrylic glass housing for safety reasons.
The separations were carried out with non-fragmented 25, 50, and 75 µm capillaries and with capillary combinations implementing a 50 µm downstream capillary segment (25 + 50, 50 + 50, and 75 + 50 µm). To find an explanation for the migration behavior also, experiments with the reverse capillary combinations were performed. In these measurements, the upstream capillary piece had an ID of 50 µm and a length of 50 cm and the downstream capillaries had IDs of 25, 50, and 75 µm and a length of 20 cm. A Lambda 1010 UV/VIS detector from Bischoff (Leonberg, Germany) was used for detection at 210 nm. The detector was placed after 40 cm. A sample solution containing 1 mmol/dm³ FcMeOH and dMFc in BGE was utilized. The injection was performed hydrodynamically by lowering the outlet buffer vial by 20 cm. A uniform sample plug was injected to compare band broadening effects. The injection segment had a length of 0.35 cm (0.5% of the total capillary length) and the respective injection times were determined based on the VFR of the corresponding capillary combination. For the electrophoretic separation, a separation voltage of 25 kV was applied and the inlet and the outlet BGE vials were placed at the same height so that there was no gravity-driven flow which affected the migration behavior. Each measurement was done three times.

**Transparent capillary connector**

To support the results from the CE-UV measurements, the behavior of a dye segment at the connection site between two capillaries with different IDs was examined utilizing a transparent capillary connector. The lab-built capillary connector was made of polyetherimide and had a 350 µm drilling. The capillary connector was additionally flattened and polished at one side for better microscope images. The lab-built capillary connector is illustrated in Fig. 6 in comparison to the commercially available capillary connector. For the experiment, a 20 cm piece of a 25 µm and a 50 cm piece of a 75 µm capillary were used. At the connection side of each capillary, 2 cm of polyimide was removed to enable visualization of the dye plug at the connection site. Both capillary segments were coupled without dead volume by means of the transparent capillary connector. The capillaries were attached with Micro Tight Nuts F 125 and with Micro Tight Sleeves F 184 from IDEX Health & Science (Oak Harbor, USA). For the images, the connector was placed on the table of a DigiMicro Lab 5.0 microscope from dnt (Dietzenbach, Germany). A saturated methylene blue solution in BGE was used as a sample solution and BGE was used as a carrier solution. The injection was performed hydrodynamically by lowering the outlet vial and capillary flow was also generated hydrodynamically. The length of the dye segment was adjusted to about 1 cm corresponding to the image section of the microscope.

**Acknowledgements** We are grateful to the German Research Foundation (DFG) for financial support (project number MA1491/12-1).

**Funding** Open Access funding enabled and organized by Projekt DEAL.

**Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

**References**

1. Ramos-Payán M, Ocaña-Gonzalez JA, Fernández-Torres RM, Llobera A, Bello-López MÁ (2018) Electrophoresis 39:111
2. Voeten RLC, Ventouri IK, Haselberg R, Somsen GW (2018) Anal Chem 90:1464
3. Beutner A, Scherer B, Matysik F-M (2018) Talanta 183:33
4. Blasco S, Kortz L, Matysik F-M (2009) Electrophoresis 30:3355
5. Beutner A, Herl T, Matysik F-M (2019) Anal Chim Acta 1057:18
6. Lapainis T, Sweedler JV (2008) J Chromatogr A 1184:144
7. Grundmann M, Matysik F-M (2012) Anal Bioanal Chem 404:1713
8. Deyl Z, Mikši­k I, Charvátová J, Eckhardt A (2003) J Chromatogr A 1013:223
9. Jooß K, Hühner J, Kiessig S, Moritz B, Neusüß C (2017) Anal Bioanal Chem 409:6057
10. Kler PA, Posch TN, Pattky M,Twigge-Jones, J, Huhn C (2013) J Chromatogr A 1297:204
11. Sydes D, Kler PA, Herrmans M, Huhn C (2016) Electrophoresis 37:3020
12. Wei J, Gu X, Wang Y, Wu Y, Yan C (2011) Electrophoresis 32:230
13. Kohl FJ, Sánchez-Hernández L, Neusüß C (2015) Electrophoresis 36:144
14. Pavlíček V, Tůma P, Matějčková J, Samcová E (2014) Electrophoresis 35:956
15. Šebestová A, Petr J (2017) Electrophoresis 38:3124
16. Tůma P, Opeká F, Samcová E (2013) Electrophoresis 34:552
17. Horká M, Karásek P, Roth M, Šlais K (2017) Electrophoresis 38:1260
18. Matysik F-M (1998) J Chromatogr A 802:349
19. Scholz R, Matysik F-M (2011) Analyst 136:1562
20. Lauer HH, Rozing GP (2018) High performance capillary electrophoresis, 2nd edn. Agilent Technologies, Germany
21. Matysik F-M, Werner G (1993) Analyst 118:1523
22. Francisco KJM, do Lago CL (2009) Electrophoresis 30:3458

Publisher’s Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.