

Maximize your CE and CEC System and experience greater laboratory uptime



Your Link to Success – Agilent Technologies Capillary Electrophoresis Capillaries, Reagents, Solutions Kits, and Supplies

Agilent Technologies, a world leader in capillary electrophoresis (CE) technology, is committed to helping you improve the quality of analyses by offering application solutions, simplifying the use of software and instrumentation, an offering unsurpassed technical support.

Agilent recognizes that CE solutions may begin with the equipment, but Agilent doesn't stop there.

Agilent provides everything you need for successful separations in CE and CEC including solutions kits, uncoated, coated and packed capillaries, reagents and buffers, supplies and spare parts. The new 2002-2003 Agilent Chromatography and Spectroscopy Supplies Reference Guide provides a complete list of all supplies to keep your lab operating at maximum efficiency.

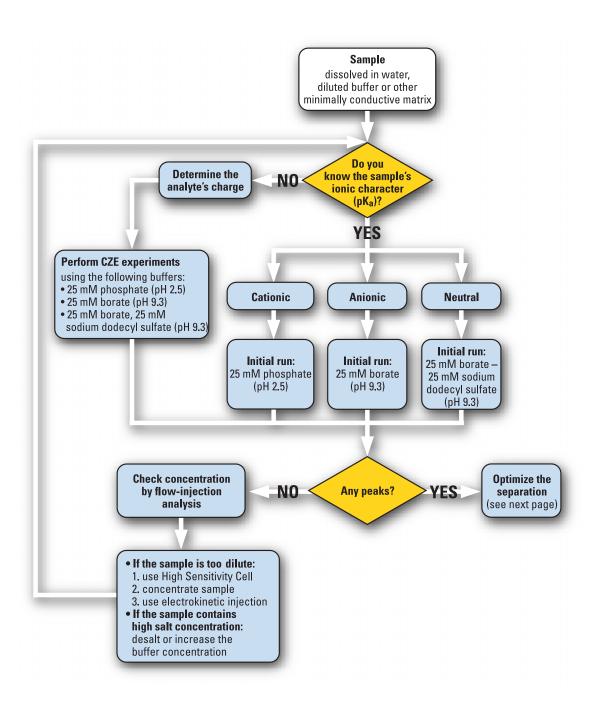


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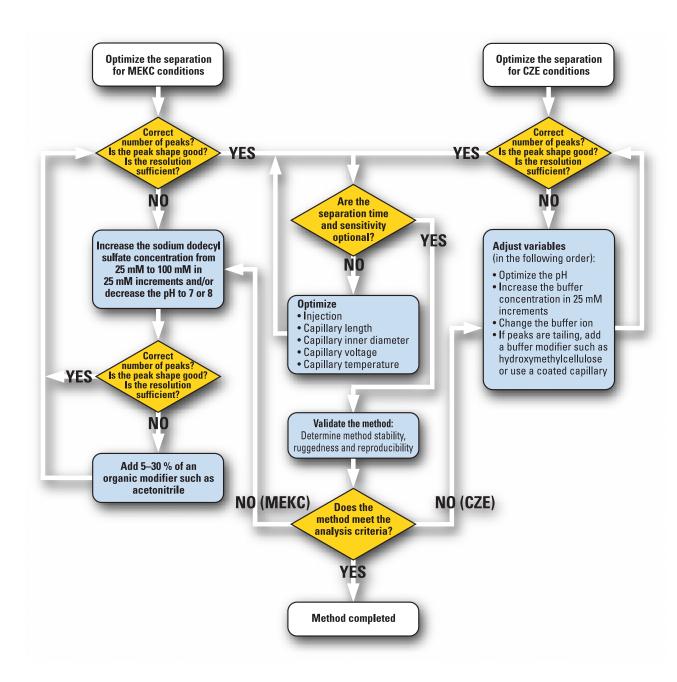
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CE Method Development



CE Method Optimization



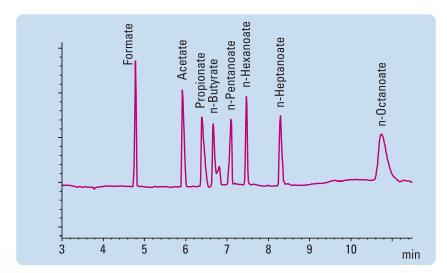
Solutions Kits for CE

Agilent continues to introduce new CE solutions kits designed to simplify many of your applications:

- plating bath solution
- forensic anions
- cations
- inorganic anions
- organic acids
- μPAGE kits for high resolution DNA fragments analysis

These kits include all you need to begin your CE analyses: buffers, capillaries, conditioning solutions, test samples, methods, and detailed descriptions. Each kit is designed to take advantage of the automation of the Agilent CE system to make your time in the laboratory more efficient. All kits are prepared using the same quality procedures as for our buffers and are thoroughly tested and supported.

While the kits have been optimized for use with the Agilent CE system, they may be used with virtually any commercial or home-built CE system.



Separation of short-chain carboxylic acids using the Organic Acids Solutions Kit



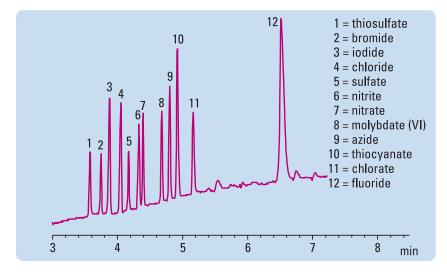
Solutions Kits for Small Ions

Inorganic Anion Solutions Kit

The Inorganic Anion Solutions Kit contains all components needed for the analysis of common inorganic anions such as chloride, bromide, iodide, fluoride, sulfate, and phosphate. Applications include the analysis of inorganic ions in:

- ultra-pure water
- waste water
- high purity chemicals
- drug formulations
- pulp and paper solutions
- semiconductor solutions

Using an indirect-UV detection system optimized for small anions, analyses are sensitive and rapid, and provide an alternative to traditional ion chromatography. The kit contains the buffer, capillaries, test mixture, and instructions.



Separation of common anions

Inorganic Anion Solutions Kit

Component	Quantity	Part Number
Inorganic Anion Solutions Kit	1	5063-6511
Buffers and Solutions		
Inorganic anion buffer	250 mL	8500-6797
Ultra pure water for CE	500 mL	5062-8578
0.1 N NaOH	250 mL	5062-8575
1.0 N NaOH	250 mL	5062-8576
Capillary		
Bare fused silica capillary, 50 µm id, L = 80.5 cm	2/pk	G1600-62211
Test Mixture		
Anion standard	10 mL	5062-8524
contains: 1000 ppm each of fluoride, chloride,		
bromide, nitrite, nitrate, sulfate; and 2000 ppm phosp	phate	
Other Components		
Product literature	1	5968-9050E

Note:

the following part should be ordered seperately when used with the Agilent CE System: Alignment Interface for standard 50 µm i.d. capillary (color code: green) Part Number: G1600-60210



Solutions Kits for Small Ions

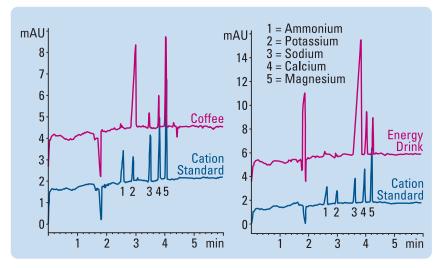
Cation Solutions Kit

The new Cation Solutions Kit provides everything you need for the analysis of inorganic and low-molecular-mass organic cations. It is specially designed for the separation of alkali metal ions, alkaline-earth metal ions and alkyl amines in a wide range of different matrices.

Each kit contains a cation buffer, bare fused silica capillaries, a cation standard, CE grade water and a detailed description of the analysis method and most common applications, including detection limits and reproducibility data. The new Cation Solutions Kit and the separation methods were developed to fit perfectly with the Agilent CE instrument and to support its high automation capabilities. The methods are easy to perform and provide accurate and quantitative analyses.

Note:

the following part should be ordered seperately when used with the Agilent CE System: Alignment Interface for 50 µm i.d. extended light path capillary, BF3 (color code: red) Part Number: G1600-60230



Cations in Coffee and Energy Drink

Food applications

- Wine Beer
- Juice Soft drinks
- Coffee
- Mineral water

Chemical/Environmental/ Pharmaceutical applications

- OTC Drugs Drinking water
- Waste water

Cation Solutions Kit

Component	Quantity	Part Number
Cation Solutions Kit	1	5064-8206
Buffers and Solutions		
Cation buffer for CE	250 mL	5064-8203
Ultra pure water for CE	500 mL	5062-8578
Capillary		
Bare fused silica capillary, extended light		
path BF3, 50 μm id, L = 64.5 cm	2/pk	G1600-61232
Test Mixture		
Cation test mixture	25 mL	5064-8205
contains: 100 ppm each Ammonium, Potassium,		
Sodium, Calcium, Magnesium		
Other Components		
Product literature	1	5968-9043E

Organic Acids Solutions Kit

The Organic Acids Solutions Kit is ideal for the analysis of short alkylchain carboxylic acids. Employing an indirect UV detection agent optimized for organic acids, the methodology is simple, sensitive, and provides accurate quantitative analysis. Suited for the analysis of organic acids in a wide range of matrices, it is especially useful for determination of organic acids in beverages, and food.

Each organic acids solutions kit contains capillaries, buffers, an organic acids standard and instructions complete with a sample electropherogram to help you begin organic acids analyses immediately.

Food applications

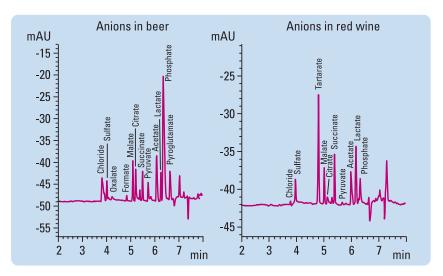
- Beer
- Wine
- Juice

Chemical applications

- Raw materials
- drug formulations

Note:

the following part should be ordered seperately when used with the Agilent CE System: Alignment Interface for 75 μ m i.d. capillary (color code: blue) Part Number: G1600-60310



Organic acids in beer and red wine

Organic Acids Solutions Kit

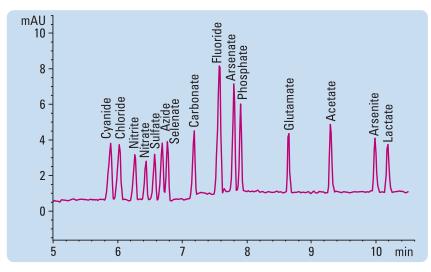
Component	Quantity	Part Number 5063-6510	
Organic Acids Solutions kit	1		
Buffers and Solutions			
Organic acids buffer	250 mL	8500-6785	
Ultra pure water for CE	500 mL	5062-8578	
1.0 N NaOH	250 mL	5062-8576	
0.1 N NaOH	250 mL	5062-8575	
Capillary			
Bare fused silica capillary, 75 μm id, L = 80.5 cm	2/pk	G1600-62311	
Test Mixture			
Organic acid test mixture	20 mL	8500-6900	
contains: 1000 ppm each, Malate, Succinate,			
Lactate			
Other Components			
Product literature	1	5968-9047E	



Solutions Kits for Small Ions

Forensic Anions Kit

This highly focused kit was developed specifically for the analysis of poisonous compounds, such as cyanide, azide, selenate, arsenate, and arsenite. In cases of poisoning, analytical tools are needed to determine the identity of the toxins quickly and accurately. A rapid determination of anionic toxins in adulterated foods and beverages is possible using CE with indirect UV detection. The forensic and other anions can be detected within 15 minutes with minimal sample preparation.



Analysis of an Anion Standard with the Forensic Anion Analysis Kit

Forensic Anions Kit

Component

Component	Quantity	Part Number
Forensic Anion Solutions Kit	1	5064-8208
Buffers and Solutions		
Basic anion buffer	5 x 50 mL	5064-8209
Ultra pure water for CE	500 mL	5062-8578
Capillary		
Bare fused silica capillary, 50 μm id, 104 cm, L = 112.5 cm	2/pk	G1600-64211
Test Mixture		
Inorganic anion test mixture	10 mL	5062-8524
containss: 1000 ppm each of fluoride,		
chloride, bromide, nitrite, nitrate, sulfate;		
and 2000 ppm phosphate		
Other Components		
Product literature	1	5968-9049E

Note:

the following part should be ordered seperately when used with the Agilent CE System: Alignment Interface for standard 50 μm i.d. capillary (color code: green) Part Number: G1600-60210



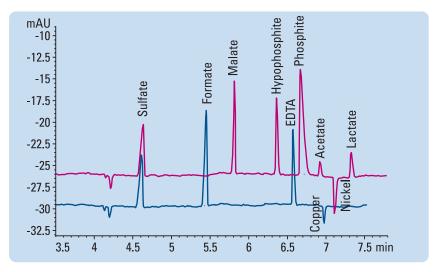
Agilent Plating Bath Analysis Kit

The Agilent Plating Bath Analysis Kit was developed for the analysis of inorganic and organic anions together with UV-absorbing metal cations (e.g. Ni²+, Cu²+, Co²+) in plating bath solutions. The bath solutions contain a variety of components such as reducing agents (which drive the plating reaction) and organic acids (as buffering and/or metal complexing agents) in addition to metal cations. Inorganic anions are also present as counter-ions of the plating metals.

This kit allows the analysis of inorganic and organic anions together with some metals in a single run. This unique feature derives from the background electrolyte's high stability constants for metal cations (which allows their analysis as anionic complexes). The kit is also suitable for the analysis of electroiron plating solutions, e.g. where different levels of Fe²⁺ and Fe³⁺ need to be monitored.

Note:

the following part should be ordered seperately when used with the Agilent CE System: Alignment Interface for standard 50 μm i.d. capillary (color code: green) Part Number: G1600-60210



Nickel- and Copper-Plating Bath

The buffer supplied in this kit is premade with the pH already adjusted so no further preparation is required. The kit also includes capillaries and a standard test mixture. To ensure that the kit and instrument are functioning properly, a detailed test procedure is provided including typical electropherograms.

Agilent Plating Bath Analysis Kit

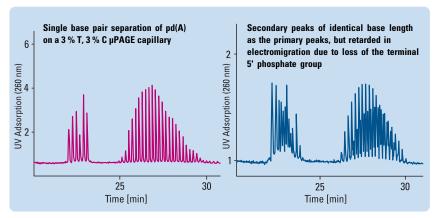
Component	Quantity	Part Number
Plating Bath Analysis Kit	1	5064-8228
Buffers and Solutions		
Plating bath analysis buffer	250 mL	5064-8236
Ultra pure water for CE	500 mL	5062-8578
Capillary Bare fused silica capillary, id = 50 µm, L = 80.5 cm	2/pk	G1600-62211
Test Mixture		
Plating bath test mixture includes: 1000 ppm each of sulfate, malate, hypophosphite, phosphite, lactate; and 611 ppm nickel	20 mL	5064-8237
Other Components		
Product literature	1	5968-9048E

μPAGE Solutions Kits for High Resolution DNA Fragments Analysis

μPAGE poly-acrylamide gel-filled capillaries are the most direct vehicles to transfer all of your applications from slab gel to CE, enjoying the automation, high speed, high resolution, and quantitative advantages of CE. The capillaries are ideal for high resolution separations of oligonucleotides, single-stranded and double-stranded DNA fragments, polymerase chain reaction (PCR) products, sequencing reaction products and oligosaccharides.

μPAGE capillaries are available in three different pore sizes. The size of the molecular sieving pores is controlled by the monomer concentration (%T) and the degree of polymer cross-linking (%C). Gels with higher %T and %C values have smaller pores and are, therefore, more effective at resolving smaller molecules. μPAGE-10 (10 %T, 0 %C) capillaries provide high resolution capabilities for separation of antisense therapeutic agents, primers and probes as well as nucleotides.





Oligonucleotide samples with or without terminal 5' phosphate group

 $\mu PAGE-5$ (5 %T, 5 %C) allows single base resolution of oligonucleotides [pd(A)] ranging from 20–150 bases while $\mu PAGE-3$ allows fast analysis of larger DNA fragments.

Three different µPAGE kits are available. For your convenience, µPAGE capillaries and µPAGE buffers can be purchased together or separately. To achieve the highest reproducibility and provide optimal longevity, use µPAGE buffer with µPAGE capillaries.

μPAGE Starter Kit

Includes 3 μ PAGE capillaries, 75 cm total length, 50 cm effective length μ PAGE pd(A)_{25-30, 40-80} oligonucleotide standard for μ PAGE-3 and μ PAGE-5 kits μ PAGE pd(A)₂₅₋₃₀ oligonucleotide standard for μ PAGE-10 kit μ PAGE buffer, 2 x 237 mL

μPAGE capillary in Kit	ID [μm]	Part Number
μPAGE-10 (10 %T, 0 %C)	100	192-1311
μPAGE-5 (5 %T, 5 %C)	75	192-5211
μPAGE-3 (3 %T, 3%C)	75	192-3211

μPAGE Basic Kit

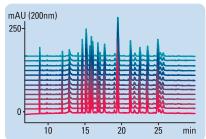
Includes 3 µPAGE capillaries, 75 cm total length, 50 cm effective length µPAGE pd(A)_{25-30, 40-60} oligonucleotide standard for µPAGE-3 and µPAGE-5 kits µPAGE pd(A)₂₅₋₃₀ oligonucleotide standard for µPAGE-10 kit µPAGE buffer, 30 mL

μPAGE capillary in Kit	ID [µm]	Part Number
μPAGE-10 (10 %T, 0 %C)	100	191-1311
μPAGE-5 (5 %T, 5 %C)	75	191-5211
μPAGE-3 (3 %T, 3%C)	75	191-3211

Standard Bare Fused-Silica Capillaries for G1600A CE Instrument

Fused-silica capillaries are the heart of CE. Capillaries from Agilent Technologies are designed and optimized for ease of use and reliability. All capillary ends are cut to a smooth, mirror-like finish using fiber optic technology. In addition, the polyimide outer-coating is removed from the ends. These processes ensure minimal sample adsorption and help maintain sharp peak shapes. All capillaries have a pre-made detection "window" and a built-in alignment stopper which allows rapid and precise insertion in the alignment interface.





CZE of a tryptic digest of recombinant human growth hormone using a standard fused silica capillary with 75 µm internal diameter

Standard Bare Fused Silica Capillaries (2/pk)

Capillary id [µm]	Total Length [cm]	Effective Length [cm]	Optical Path Length [µm]	Color code	Part Number
50	33	24.5	50	•	G1600-63211
	48.5	40	50		G1600-60211
	64.5	56	50		G1600-61211
	80.5	72	50		G1600-62211
	112.5	104	50		G1600-64211
75	33	24.5	75		G1600-63311
	48.5	40	75		G1600-60311
	64.5	56	75		G1600-61311
	80.5	72	75		G1600-62311
	112.5	104	75		G1600-64311
100	33	24.5	100		G1600-63411
	48.5	40	100		G1600-60411
	64.5	56	100		G1600-61411
	80.5	72	100		G1600-62411
	112.5	104	100		G1600-64411

Note:

All capillaries are supplied in package size of 2/pack.

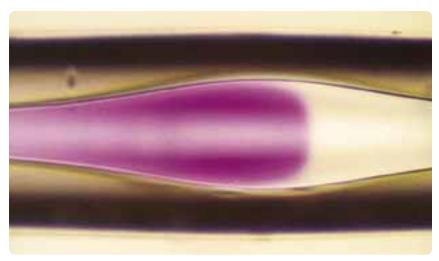
Helpful hint:

Different inner diameters of capillaries need to use different alignment interfaces to guarantee optimal detection. The color coding of the capillary and the alignment interface allow you to easily match the correct interface with the capillary.

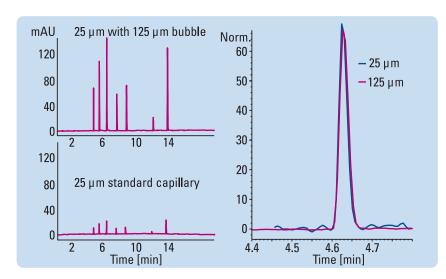
Extended Light Path Capillaries (Bubble Cell Capillaries) Bare Fused-Silica

Use Agilent Technologies Extended Light Path Capillaries ("bubble" cell capillaries) to improve sensitivity 3-to 5-fold over standard capillaries. With Extended Light Path Capillaries the inner diameter is increased only at the detection window, offering the sensitivity of a wide inner diameter capillary and the low current generation of a narrow one.

Resolution is not sacrificed when used with Agilent Technologies' matching optical alignment interfaces.



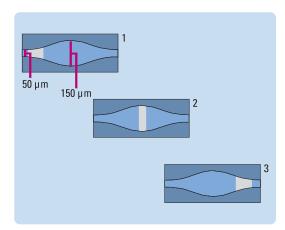
Enhanced sensitivity with Agilent Extended Light Path Capillaries ("bubble" cell)



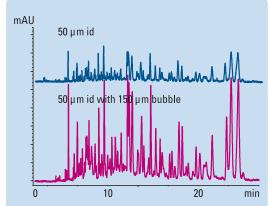
Analysis of cold medicine ingredients in a standard capillary (id 25 µm) and an Agilent Extended Light Path Capillary Through a computer-controlled, proprietary process the diameter is increased three to five times with a manufacturing precision better than 3%. Take advantage of this process to extend the detection pathlength of 25 μ m id capillaries to 125 μ m, 50 μ m to 150 μ m, and 75 μ m to 200 μ m!

Helpful Hint:

Use narrow 25 and 50 µm id "bubble" cell capillaries for highly conductive buffers without sacrificing sensitivity



Electroosmotic flow maintains the "plug" flow in the bubble. Optical slits matched to the zone geometry maintain resolution



CZE analysis of a tryptic digest of carbonic anhydrase using a standard capillary (id 50 $\mu m)$ compared with an Agilent Extended Light Path Capillary

Agilent Technologies Extended Light Path Capillaries Bare Fused-Silica (2/pk)

Capillary id [µm]	Total Length [cm]	Effective Length [cm]	Bubble Factor	Optical Path Length [µm]	Color Code	Part Number
25	48.5	40	5	125	•	G1600-60132
	64.5	56	5	125	•	G1600-61132
	80.5	72	5	125	•	G1600-62132
50	48.5	40	3	150	•	G1600-60232
	64.5	56	3	150		G1600-61232
	80.5	72	3	150		G1600-62232
	112.5	104	3	150		G1600-64232
75	48.5	40	2.7	200	•	G1600-60332
	64.5	56	2.7	200		G1600-61332
	80.5	72	2.7	200		G1600-62332
	112.5	104	2.7	200		G1600-64332

For more information refer to: D. N. Heiger, P. Kaltenbach, H-J. P. Sievert, Electrophoresis, 1994, 15(10), 1234-1247 and PN 5963-1889E.

Universal Bare Fused-Silica Capillaries

These are capillaries with a window and 75 cm effective length and, 363 μm od, fitting into any CE instrument. To cut them to the correct length we recommend using the P/N 5183-4669 CE column cutter.

Capillary id [µm]	Total Length [cm]	Effective Length [cm]	Part Number
20	100	75	190-0431
50	100	75	190-0131
75	100	75	190-0231
100	100	75	190-0331

Bulk Fused-Silica Capillaries, 363 µm od

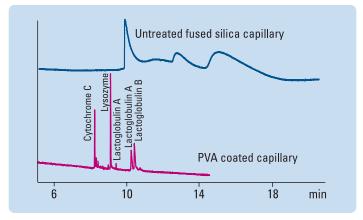
Capillary id [µm]	Total Length [m]	Part Number	
20	5	160-2660-5	
50	5	160-2650-5	
75	5	160-2644-5	

Coated Capillaries

Polyvinyl Alcohol-Coated (PVA) Capillaries

PVA capillaries contain a permanently adsorbed layer of polyvinyl alcohol. This coating shields the silanol groups of the fused silica and effectively eliminates electroosmotic flow (EOF). Using a proprietary deposition process, the PVA coating is stable over a wide pH range - even under basic conditions - from pH 2.5 to 9.5. This stability allows use of a wide range of common CE buffers. Because the silica surface is covered, many proteins and amines can be analyzed without the peak tailing found with uncoated capillaries. In addition, since EOF is eliminated, cumbersome washing procedures are unnecessary and migration time reproducibility can be improved.

PVA capillaries can be used for a variety of applications, including protein analysis at physiological pH, isoelectric focusing, and small anion analysis without the need for flow-reversal agents in the buffer.



Use of PVA capillaries to reduce protein adsorption



91 - 2-Amino-5-azotoluene uim - 2-Amino-5-azotol

CZE analysis of basic amines using PVA capillaries (decomposition products of azo dyes)

For Agilent Capillary Electrophoresis System Users

Capillary id [µm]	Total Length [cm]	Effective Length [cm]	Bubble Factor	Optical Path Length [µm]	Color Code	Part Number	Comment
50	64.5	56	none	50		G1600-61219	standard
	64.5	56	3	150		G1600-61239	extended light path
50	125	21.6	none	50	•	G1600-67219	for CE-MS
75	64.5	56	none	1200		G1600-68319	for HSDC
	125	21.6	none	75	•	G1600-67319	for CE-MS
100	48.5	40	none	100		G1600-60419	standard
	64.5	56	none	100		G1600-61419	standard

Note: The PVA capillaries for CE-MS have a blue alignment stopper, matching the blue color code of the alignment interface for MS-UV-Vis detection. The alignment stopper of the 50 µm id PVA capillary for CE-MS has a black dot for easy identification.

For non-Agilent CE System Users

Capillary id [µm]	Total Length [cm]	Effective Length [cm]	Bubble Factor	Optical Path Length [µm]	Part Number
50	71	60	none	50	G160U-61219
	71	60	3	150	G160U-61239
100	56	45	none	100	G160U-60419
	71	60	none	100	G160U-61419

Note:

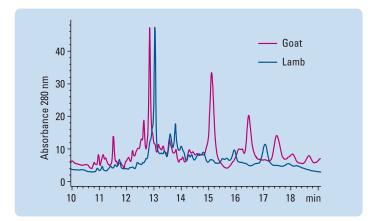
When extended pathlength capillaries are used in non-Agilent systems, loss of resolution may be found if the axial slit width is not reduced. In Agilent Technologies systems the alignment interface contains properly matched slits to maintain resolution (for details, see PN 5963-1889E).

greater lengths for use in non-Agilent systems! Now the PVA is also available for use

The PVA coating is available in standard capillaries, or in Agilent Extended Light Path Capillaries ("bubble" cell capillaries) for high sensitivity applications. Both

capillary types are available in

with the High Sensitivity Detection Cell for even further improved HPLClike sensitivity. In addition PVA coated capillaries are offered for CE-MS applications. The capillaries are provided with normally positioned detection window to allow tandem UV-Vis and MS detection for improved sample identification.



Analysis of meat proteins by c-IEF using PVA capillaries

Poly(vinyl alcohol)-coated Capillaries - A Solution for many Application Areas on Capillary Electrophoresis U. Jegle, T. Soga, R. Grimm, H. Godel, R. Schuster and G. Ross, LC-GC International, March 1997.

For more information refer to:

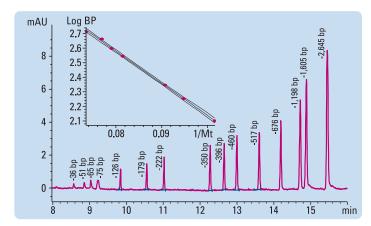
Coated Capillaries

CEP Coated Capillaries

CEP capillaries contain a permanently bonded polymer coating. This CEP coating shields the silanol functionality of the capillary surface and helps prevent sample adsorption. Additionally, the EOF is nearly eliminated making the capillary ideal for applications such as DNA separations with sieving polymer buffers (see dsDNA kit, page 13). Elimination of EOF also simplifies analysis of anions and organic acids by direct UV detection. Without EOFreduction, highly mobile ions such as nitrate can migrate in the opposite direction to the slower, longer chain acids.

The CEP coated capillary is stable from pH 2 to 8. It can be used with borate buffers offering a different surface functionality to help alleviate sample adsorption.

Each batch of CEP coated capillaries is rigidly tested by Agilent Technologies and includes a representative electropherogram to assure quality.



Restriction fragment separation (36-2645 bp)

Capillary id	Total	Effective	Optical	Part Number
[µm]	Length [cm]	Length [cm]	Path Length [µm]	
75	97	72	75	G1600-62318

For Agilent CE systems and non-Agilent CE systems

Bulk µSIL-DB Capillaries

The μ SIL-DB coated capillaries are available as μ SIL-DB-1 and μ SIL-DB-17. In combination with a cellulose based buffer system, μ SIL-DB coated capillaries have been widely used in cIEF applications, PCR product and DNA fragment separation and many other CE applications which require reduced EOF.

	FIIII						
Capillary	Capillary ID [µm]	Length [m]	Thickness [µm]	Part Number			
μSIL-DB-1	50	10	0.05	126-1012			
μSIL-DB-1	50	10	0.20	126-1013			
μSIL-DB-1	100	10	0.10	127-1012			
μSIL-DB-17	50	10	0.10	126-1713			
μSIL-DB-17	100	10	0.10	127-1712			
μSIL-DB-17	100	10	0.20	127-1713			

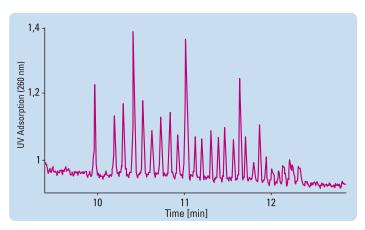
Cross-linked and Bonded µSIL Capillaries

μSIL-FC and μSIL-DNA Capillaries With Windows

A series of coated capillaries specifically designed for CE which are prepared by cross-libking and bonding a novel, proprietary fluorocarbon (FC) polymer. µSIL-FC capillaries are chemically inert, hydrophobic, and stable from pH 2.5—10.0.

These capillaries are a must-have for cIEF, protein, peptide and carbohydrate separations, as well as replaceable gel CE applications such as oligonucleotides, DNA fragments, and PCR product separations.

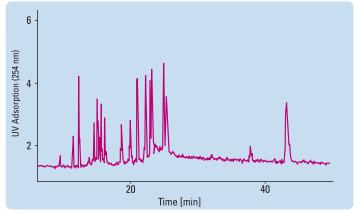
 $\mu SIL\text{-}DNA$ capillaries are also coated with an FC polymer but have a 75 μm id to accomodate the viscosity of entangled polymer solutions. All μSIL capillaries are batch tested to ensure the highest performance and reproducibility.



Analysis of Allelic ladder with µSIL-DNA

μSIL-WAX Capillaries With Windows

µSIL-WAX features a modified, polyethylene oxide, hydrophilic coating made through a special cross-linking and bonding process. The coating effectively masks active silanol sites, offering exceptional efficiency, resolution, peak shape and reproducibility. The highly stable coating and near-zero EOF of μ SIL-WAX makes the capillary ideal for CE-MS, and protein and peptide separations from pH 2–5.



Analysis of Myoglobin tryptic digest using $\mu SIL\text{-WAX}$

Capillary	Capillary ID [µm]	Total Length [cm]	Effective Length [cm]	Film Thickness [µm]	Unit	Part No.
μSIL-FC μSIL-DNA	50 75	80 65	50 50	0.075 0.075	3/pk 2/pk	194-8111 199-2602
μSIL-WAX	50	100	75	0.1	2/pk	196/7203
μSIL-WAX	100	100	75	0.1	2/pk	197-7202



Capillaries for Capillary Electrochromatography (CEC)

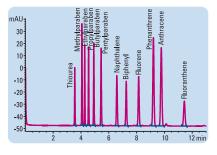
Capillary Electrochromatography is a hybrid of CE and LC and can be performed in the Agilent CE system. Using CE capillaries packed with LC stationary phases, CEC offers the loadability and selectivity of LC and the high efficiency of CE.

Using the high pressure capabilities of the Agilent CE system, both ends of the CEC capillary can be pressurized. This process prevents outgassing upon application of high voltage and significantly extends capillary lifetime.

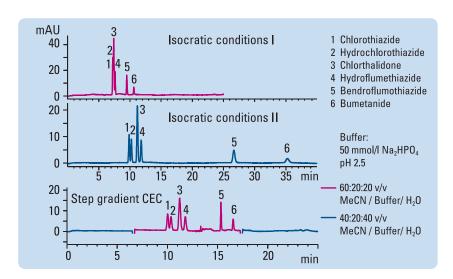
Use CEC to improve resolution of solutes which are difficult to resolve by HPLC, for hydrophobic solutes which cannot be solubilized in MEKC buffers, or for reduced sample and solvent consumption compared to HPLC.

Helpful Hint:

CEC capillaries require an Agilent CE system with external gas supply capabilities.



Capillary Electrochromatography of parabenes and aromatics



Capillary Electrochromatography of diuretic test mixture (courtesy of Dr. Melvin Euerby, Astra Charnwood, UK)

Standard CEC Capillaries

Description		Dimensions length × id [mm]	Detection	Part Number
CEC-Hypersil C18,	3 µm	250 × 0.1	Standard	5063-6512
CEC-Hypersil C18,	3 µm	400 × 0.1	Standard	5063-6513
CEC-Hypersil C8,	3 µm	250 × 0.1	Standard	5063-6535
CEC-Hypersil C8,	3 µm	400 × 0.1	Standard	5063-6540
CEC-Hypersil Phenyl,	3 μm	250 × 0.1	Standard	5063-6536
CEC-Hypersil Phenyl,	3 μm	400 × 0.1	Standard	5063-6541

CEC Capillaries for HSDC*

Description		Dimensions length × id [mm]	Detection	Part Number
CEC-Hypersil C18,	3 µm	250 × 0.1	HSDC*	5063-6537
CEC-Hypersil C18,	3 µm	400 × 0.1	HSDC*	5063-6542
CEC-Hypersil C8,	3 μm	250 × 0.1	HSDC*	5063-6538
CEC-Hypersil C8,	3 μm	400 × 0.1	HSDC*	5063-6543
CEC-Hypersil Phenyl,	3 µm	250 × 0.1	HSDC*	5063-6539
CEC-Hypersil Phenyl,	3 μm	400 × 0.1	HSDC*	5063-6544

^{*}use with High Sensitivity Detection Cell (HSDC) only

For test sample order: 01080-68704

Alignment Interfaces and Capillary Cassette

Agilent Technologies alignment interfaces are an integral part of the Agilent diode-array detection system. These interfaces contain optical slits which are precisely matched to the capillary inner diameter for optimized sensitivity and linear detection range.



Alignment Interfaces

Description	Color Code	Corresp. capillary	Part Number
Alignment interface for standard capillaries with id 50 µm			G1600-60210
Alignment interface for standard capillaries with id 75 and 100 µm	•		G1600-60310
Alignment interface for Agilent Extended Light Path capillaries with id 25 μm	•	•	G1600-60150
Alignment interface for Agilent Extended Light Path capillaries with id 50 µm		•	G1600-60230
Alignment interface for Agilent Extended Light Path capillaries with id 75 µm		•	G1600-60330
PEEK alignment interface for CE-MS (360 od capillaries)			G1600-60400

Notes

- 1. Standard 75, and 100 μm id capillaries use the same interface (blue).
- 2. PVA coated 50 and 75 μm id capillaries for CE-MS use the same non-metallic interface with color code green.

Cassette

Description	Part Number
Capillary cassette for use with standard capillaries, extended	
light path capillaries and the High Sensitivity Detection Cell	G1600-60002

Helpful Hint:

The cassette and interfaces accept all commercially available capillaries (\sim 365 μ m od).

In combination with the capillary cassette, alignment interfaces simplify capillary exchange, protect the fragile detection window and ensure exact alignment of the window in the detector.



Agilent Technologies High Sensitivity Detection Cell

The Agilent High Sensitivity Detection Cell Dramatically Enhances Sensitivity and Linearity for the Agilent CE Capillary Electrophoresis System

The Agilent High Sensitivity

Detection Cell — a technological leap which extends sensitivity by an order of magnitude — provides a solution to sensitivity limitations often encounter in CE. This improvement will substantially increase the utility of CE for impurity analysis of chiral drugs, biologicals, and compounds of environmental interest, among others.

The High Sensitivity Detection Cell for the Agilent CE system not only improves detection sensitivity more than 10-fold over standard capillaries, but also extends linearity beyond 2000 mAU and provides unsurpassed spectral fidelity. These improvements are a result of a proprietary micromachined design which increases the detection pathlength from 75 µm to 1200 µm while dramatically reducing stray light.

The High Sensitivity Detection Cell has a design comprised of a fused silica cell body and removable capillaries. The light path through the cell is made from black fused silica which significantly minimizes stray light and defines the aperture for the diode-array spectrometer. In addition, the reflective interior functions as a "lightpipe", insuring almost 100 % transmission of light which entered the cell. These properties result in enhanced linearity and unsurpassed spectral fidelity with the diode-array detector.

The high linear range allows quantification of both < 0.1% impurities and the main component in one run. This is useful for all impurity analyses and is especially useful for determining chiral excess.

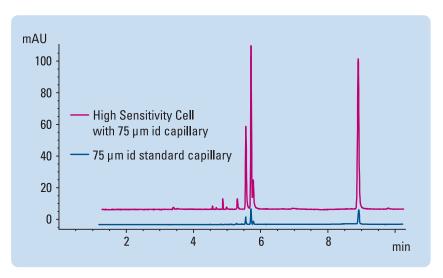
Integral to the Agilent High Sensitivity Detection Cell function is maintenance of peak shape. The unique geometry of the capillaries insure proper coupling with the cell, virtually eliminate dead volume and maintain peak shape. The end of these capillaries include both a flaring of the inner diameter and beveling of the outer diameter.

The Agilent High Sensitivity
Detection Cell can be used in all
Agilent CE systems, regardless of
age. As shown in the figure the cell
is incorporated into an optical
interface which is similar to that
used for standard and bubble cell

capillaries – making capillary and cassette installation the same as for the other capillaries. The figure also illustrates the decoupled design which allows quick and easy replacement of capillaries. This process is simplified by use of preassembled ferules and finger-tight fittings.







Agilent High Sensitivity Detection Cell vs. 75 µm Standard Capillary for the CZE separation of naphthalene sulfonic acids

Characteristics of the Agilent High Sensitivity Detection Cell

- 10-fold increase in signal-to-noise
- Detector linearity beyond 2000 mAU for accurate quantitative analysis
- Decoupled design allows replaceable capillaries and reduced cost operation
- Special capillary geometry insures maintenance of peak symmetry
- Full diode-array spectral capabilities
- Design fits all Agilent CE instruments

High Sensitivity Detection Cell (HSDC)

Description	Part Number
High Sensitivity Detection Cell Kit	G1600-68713
Detection cell, 75 µm id inlet capillary (72 cm) and outlet	
capillary (8.5 cm) pair, capillary cassette, fittings (3 fitting	
screws with seals, 2 fitting caps), 4.5 mL cleaning solution	
Replacement Capillary Kits: Bare Fused Silica	
75 µm id capillary pair with 8.5 cm outlet and the following inlet:	
56 cm effective length	G1600-68716
72 cm effective length	G1600-68715
88 cm effective length	G1600-68714
Replacement Capillary Kit: PVA Coated Capillaries	
75 µm id capillary pair with 8.5 cm outlet and the following inlet:	
56 cm effective length	G1600-68319
Replacement Fittings	G1600-63200
3 fitting screws with seals, 2 fitting caps	
Replacement Detection Cell	G1600-60027
Cleaning Solution 1.3 kg	5062-8529

Notes

- A modified capillary cassette is required for the Agilent High Sensitivity Detection Cell.
 This cassette is included with the kit and can also be used for all capillaries and optical interfaces.
- 2. CEC capillaries are also available for the HSDC (for part numbers see page 19).

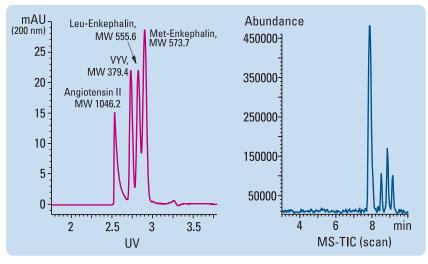
For more information refer to: Publication No. 5965-5984E.

CF-MS Accessories

The G1603A CE-MS Adapter kit includes a MS-cassette, two bare fused silica capillaries and the accessories to prepare the Agilent CE to be coupled to the electrospray mass spectrometer. It does not contain the accessories needed to prepare the MS. The CE-MS adapter kit can be used with the Agilent 1100 Series MSD, and MSD Trap System or virtually any other electrospray MS-platform.

The CE-MS cassette completely thermostats the capillary until it exits the CE system. A methods development configuration uses online diode array detection (DAD) and MS. For rapid or routine MS analysis the DAD can be by-passed to decrease the total capillary length and reduce analysis time.

The CE-ESI-MS Nebulizer Kit includes the electrospray needle and splitter assembly which allows the direct connection of the CE-instrument with the Agilent 1100 Series MSD. The CE-ESI-MS



CE-MS of 4 component peptide mixture (210 fmol)

Bebulizer Kit needs the CE-MS Adapter Kit for fully supporting the CE-MS coupling.

The CE with tandem UV-Vis and MS detection allows the analysis of complex mixtures. Analyte mixtures are separated and the components detected via UV-Vis absorption,

allowing preliminary identification based on peak elution time and/or UV-Vis spectra when compared to a standard. The on-line coupling to electrospray-ionization mass spectrometry (ESI-MS) then reveals unambiguous information on the solute's molecular weight and possibly also structure.



Agilent CE-MS Adapter Kit (1)

Description	Part Number	
Includes CE-MS interface cassette, non-metallic alignment interface for 360 µm od capillaries, CE-MS capillary (id 50 µm, I=21.6 cm, L=125 cm), protective tubing and warning label, ground cable	G1603A	
The following parts of the kit can be ordered separately:		
CE-MS cassette, metallic	G1600-60013	
PEEK alignment interface for 360 μm od capillaries	G1600-60400	
CE-MS capillary, bare fused silica, id 50 µm, l=21.6 cm, L=125 cm (2/pk)	G1600-67311	

(1) Interfacing the capillary requires an electospray needle which is not included in this kit but in the CE-ESI-MS Nebulizer Kit. For coupling with non-Agilent MS please contact the MS vendor.

CE-ESI-MS Nebulizer Kit for 1100 LC/MSD and LC/MSD Trap

Quantity	Part Number
	G1607A
ordered separately:	
1	G1607-60041
1	G1607-60034
1	8710-2164
1	5022-2142
1	G1607-60001
1	G1607-60000
1	5022-2141
2	0100-1543
2	1520-0401
2	0515-0982
2	G1607-20030
2	0100-1544
1	8500-4410
1	5063-6590
	e ordered separately: 1 1 1 1 1 1 1 2 2 2 2 2 2 1

Capillaries for CE-MS

Description	Color Code	Part Number
CE-MS capillary, bare fused silica, i.d. 50 µm, l=21.6 cm, L=125 cm (2/pk)	•	G1600-67311
CE-MS capillary, PVA coated, i.d. 50 µm, l=21.6 cm, L=125 cm (1/pk)	•	G1600-67219
CE-MS capillary, PVA coated, i.d. 75 µm, I=21.6 cm, L=125 cm (1/pk)	•	G1600-67319

For more detailed information refer to the technical note:

CE-ESI-MS: An Integrated Solution – PN 5968-1328E

Buffers and Reagents for CE

The premade buffers, ready-to-use right out of the bottle, help eliminate the time-consuming buffer preparation process.

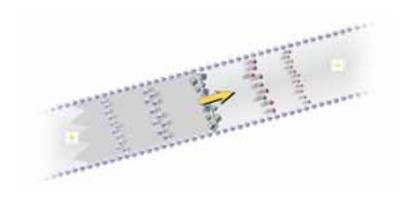
In addition to a set of kit buffers, which are specially designed for dedicated applications, Agilent offers a series of basic CZE buffers covering a broad pH range. Further the product portfolio includes special buffers for protein analysis and for Micellar Electrokinetic Chromatography (MEKC). Cleaning and conditioning solutions complete the offer.

All Agilent buffers and reagents are designed to meet the stringent demands of CE. Manufactured under GLP/GMP conditions in ISO9001 facilities, each is shipped with assay information and verification of purity. Chemicals are all electrophoresis grade, with nearly all ionic and organic impurities removed. Solutions are prepared under Class 10 cleanroom conditions and prefiltered through 0.2 mm filters to ensure removal of particulates.

To maintain high reproducibility all buffer solutions are standardized to 0.02 pH units in accredited laboratories. Superior quality control ensures reproducible results — bottle-to-bottle and batch-to-batch.

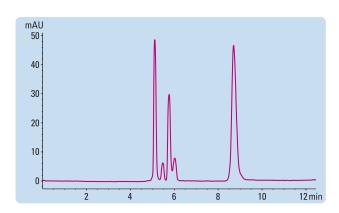






Int. (EmL/Ref) 150 200 Ex [nm] Em [nm]

The total fluorimetry spectrum of the 50 mM borate buffer pH 9.2 verifies that the solution is free of fluorescence-active impurities (1 and 2 = Rayleigh stray light of zero and first order, 3 = Raman stray light)



CZE analysis of a peptide mixture using premade 50 mM sodium phosphate buffer, pH 2.5 $\,$

CE-grade Water

Item	Volume	Part Number
Ultra pure water for CE	500 mL	5062-8578

Capillary Conditioning Solutions

Item	Volume	Part Number
0.1 N sodium hydroxide solution	250 mL	5062-8575
1.0 N sodium hydroxide solution	250 mL	5062-8576
0.1 N phosphoric acid solution	250 mL	5062-8577

CZE Buffers for Charged Analytes

Item	Volume	Part Number
50 mM sodium phosphate buffer, pH 2.5	250 mL	5062-8571
50 mM sodium phosphate buffer, pH 7.0	250 mL	5062-8572
50 mM sodium tetraborate buffer, pH 9.3	250 mL	5062-8573
20 mM sodium tetraborate buffer, pH 9.3	100 mL	8500-6782

Buffers for CZE of Proteins

Item	Volume	Part Number
50 mM phosphate, 0.05% hydroxyethyl	250 mL	8500-6786
cellulose buffer, pH 2.5		
150 mM phosphate, 200 mM ammonium	250 mL	8500-6787
sulfate buffer, pH 7.0 1)		

 $^{^{1)}\,\}mbox{Use}$ 25 $\mu\mbox{m}$ id capillary to limit current generation (see page 14)

Micellar Electrokinetic Chromatography (MEKC) Buffer for Neutral and Charged Analytes

Item	Volume	Part Number
50 mM sodium tetraborate, 100 mM sodium dodecyl sulfate buffer, pH 9.3	250 mL	5062-8574

Dilute with 50 mM sodium tetraborate, pH 9.3 (PN 5062-8573) to reduce SDS concentration without affecting the tetraborate composition or pH $\,$

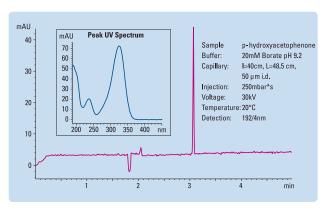
μPAGE Buffer Solutions and Oligo Standards

Agilent CE System Start-up and Test Kits

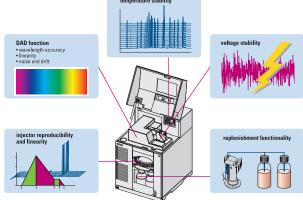
Agilent Technologies offers chemical test kits and validation packages to help comply with regulatory and quality standards. The Installation Qualification (IQ) Chemical Kit and Hardware Start-up Kit which are shipped with new instruments are useful for rapidly

verifying system functionality. For rigorous testing, the Operational Qualification (OQ) / Performance Verification (PV) Kit can be used to verify DAD noise, drift, linearity, and wavelength accuracy, and replenishment functionality.

The OQ/PV Kit is only part of the validation services available from Agilent Technologies. When implemented by our qualified personnel it can be used to help validate your Agilent CE system.



IQ and OQ/PV test method



Operational Qualification / Performance Verification - 0Q/PV

Agilent CE Hardware Start-up Kit

Item	Part Number
Includes one Extended Light Path capillary (I 56 cm, id 50 µm),	G1600-68706
one standard capillary (I 56 cm, id 50 μm),	
one test capillary (I 40 cm, id 50 μm),	
one alignment interface for standard capillaries (id 50 µm),	
and one alignment interface for Extended Light Path capillaries (id.	50 um)

r^2 = 0.9992 1 0 200 400 600 Injection (mbar*s)

OQ/PV injection linearity test

Agilent CE Installation Qualification (IQ) Chemical Kit

item	Part Number
Includes buffer 20 mM borate, pH 9.2, 100 mL,	5063-6514
test sample 4-(hydroxy)acetophenone 1 mM, 2 mL,	
capillary conditioning solution 0.1 N sodium hydroxide, 100 mL,	
and manual with methods and instructions	

Agilent CE Operational Qualification (OQ) / Performance Verification (PV) Kit

Item	Part Number
Includes buffer 20 mM borate, pH 9.2, 100 mL,	5063-6515
test samples 0.1, 0.5, 1.0, and 5.0 mM 4-(hydroxy)- acetophenone, 2 mL ea.,	
capillary conditioning solution 0.1 N sodium hydroxide, 100 mL,	
test capillary I = 40 cm, L = 48.5 cm, id 50 μm,	
diskette with methods, sequence, and spectral library, and instruction manual	
00/PV Chemical Kit includes buffer and test sample	5063-6520

Capillary Electrophoresis Supplies and Accessories







Vials and Caps

Item	Part Number
Buffer vials	
Vial polypropylene, 1 mL (100/pk)	5182-0567
Buffer vials for CE, clear glass, 2 mL (100/pk)	5182-9697
Buffer vials for CE, clear glass, 2 mL (500/pk)	5183-4623
Buffer vials for CE, amber glass, 2 mL (100/pk)	5183-4619
Buffer vials for CE, amber glass, 2 mL (500/pk)	5183-4622
Sample vials	
Vial polypropylene, 100 μL (1000/pk)	9301-0978
Vial glass-lined polypropylene, 100 mL (100/pk)	9301-0977
recommended for routine use	
Standard vial caps	
Vial caps PUR (polyurethane for re-sealing) (100/pk)	5181-1512
Vial caps PEO (polyethylene olefin for chemical resistance) (100/pk)	5181-1507
Vial caps PEO (polyethylene olefin for chemical resistance) (500/pk)	5181-1513

Accessories

Description	Part Number
50-vial rack (5/pk)	9301-0722
Ceramic capillary scribes (4/pk)	5181-8836
CE capillary column cutter	5183-4669
Diamond blade replacement kit for CE cutter	5183-4670
Windows etching tool, 3/pk	590-3003

Agilent CE System Accessory Kit



Item	Part Number
Includes electrode tool, screwdriver, fuses, air filter, glass frit,	G1600-68701
PP vials and PUR caps	

Spare Parts

Item	Part Number
Deuterium lamp	2140-0585
Electrode assembly, standard	G1600-60007
Electrode assembly, short	G1600-60033
12 mm socket wrench (electrode tool)	8710-2076
O-ring silicone (5/pk)	5062-8544
Pre-puncher	G1600-67201
Electrolyte bottle	9300-1748
Glass frit	5041-2168
Filter frit adapter	5062-8517
Bottle cap for electrolyte bottle	9300-1747
Bottle sealing 0-Ring	0905-1163
Plug for electrolyte bottle	G1600-23223
Air filter	3150-0619
Plastic screw box for puncher, insulation plate (10/pk)	G1600-62402
260 nm diode-array filter for DNA analysis with polyacrylamide-filled capillaries and oligonucleotide analysis	G1600-62700

Agilent Technologies: Quality CE Support and Applications Literature

Agilent Technologie's involvement in CE was inspired by the initial ground work of Professor J. W. Jorgenson in 1981. Research at Agilent began in the early 1980s with a series of fundamental publications by McManigill and coworkers at Agilent laboratories.

Since then Agilent Technologies has created a network of specialists all over the world. With major research groups in the US and Germany, application experts in the US, Germany, and Japan, and CE field specialists all over the world, Agilent is ready to help you with your separation challenges. A continuous stream of publications from Agilent, ranging from educational primers to scientific publications in respected journals, helps keep you up to date with the latest developments and trends in Capillary Electrophoresis.

Please contact Agilent
Technologies for your free-ofcharge copy of the literature listed
on these pages, or visit us on the
world wide web.



A Selection of Agilent Technologies Capillary Electrophoresis Application Literature

Application Notes

Title	Publication N
Double-stranded DNA Analysis with the Agilent Capillary Electrophoresis System	5988-4304EN
Dligonucleotide Analysis with the Agilent Capillary Electrophoresis System	5988-4303EN
Analysis of human rhinovirus (common cold virus) in viral preparations by CZE	5988-2591EN
Analysis of paraquat and diquat by CE/MS	5988-1664EN
Analysis of peptides using CE/MS/MS	5988-1426EN
CE and CE/MS for the Analysis of Natural Products	5980-2081E
Analysis of fangchinoline and tetrandrine in Chinese traditional medicines by capillary electrophoresis	5980-0456E
Quantitation of the alkaloids berberine, palmatine and jatrorrhizine in the Mahonia stem	5980-0455E
Analysis of chlorogenic acid in traditional Chinese medicines by capillary electrophoresis	5980-0457E
Rapid Screening of Amino Acids in Food by CE-ESI-MS	5968-8952E
Optimized parameters for the analysis of tropane alkaloids by CE-ESI-MS	5968-9414E
Analysis of Drugs of Abuse by CE-ESI-MS	5968-9221E
Analysis of Hallucinogenic Mushrooms by CE-ESI-MS	5968-9219E
Frace Anion Determination in Semiconductor Grade by Capillary Electrophoresis	5968-8953E
ASTM approved method for water analysis by capillary electrophoresis	5968-8660E
Simultaneous analysis of inorganic anions, organic acids, amino acids and carbohydrates using the Basic Anion Buffer	5968-7715E
Analysis of Aromatic Amines in Leather Extracts by CE-MS	5968-7929E
Rapid Monitoring of Carbohydrates in Food with Capillary Electrophoresis	5968-6985E
Analysis of Mahuang Chinese herbal medicine by capillary electrophoresis	5968-6418E
Analysis of Poisoned Food by Capillary Electrophoresis	5968-5731E
Monitoring of Electroless Plating Baths by Capillary Electrophoresis	5968-5761E
Analysis of Flavonoids in Plant Extracts by CE-MS	5968-5729E
Analysis of Amphetamines in Urine by CE-ESI-MS	5968-3879E
Analysis of Sulfur Anions in Kraft Liquors Using Capillary Electrophoresis	5968-3306E
lighly Sulphonated Cyclodextrins for Chiral Analysis	5968-2018E
Analysis of Anionic Contamination on Wafer Surfaces of Semiconductors	5966-4601E
Development of a method for separation of the four sterioisomers of troglitazone using capillary electrophoresis	5966-3111E
Analysis of Anions in Power Plant Waters using Capillary Electrophoresis	5966-3741E
Second derivative spectral identification of tryptophan and tyrosine in peptides	5966-2958E
Purity check of a synthetic alpha-homo polylysine preparation	5966-2956E
Capillary zone electrophoresis of neuropeptides	5966-2955E
Micropreparative capillary zone electrophoresis – tryptic digest analysis of recombinant Gr	oES 5966-2947E
Digonucleotides—Fraction Collection from Capillary Gel Electrophoresis and Offline MALDI-	TOF 5965-9036E
Digonucleotide Analysis by Capillary Gel Electrophoresis	5965-9037E
Digonucleotides Analysis of Antisense Therapeutics	5965-9038E
ligh Sensitivity SDS-protein separations by Capillary Electrophoresis	5965-9035E
Iltra-Low Level Impurity Analysis by Capillary Zone Electrophoresis	5965-9034E
ligh Sensitivity Chiral Excess Analysis	5965-9033E
Fransfer of HPLC methods to Capillary Electrochromatography	5965-9031E
Gradient LC analysis of herbicides and polyaromatic hydrocarbons by isocratic Capillary Electrochromatography	5965-9032E

Publication No
5965-9030E
5965-9028E
5964-0280E
5963-7568E
5963-1190E
5963-1122E
5091-9062E

In addition to the application notes
Agilent provides so-called application
briefs. An application brief describes
the separation conditions for a wide
range of different samples in a
comprehensive format. For an overview
and the content of the Agilent
application briefs visit our Agilent
home page on the worldwide web:
http://www.agilent.com/chem

Scientific Journal Reprints

Title	Publication No.
CE/MS: Practical Implementation and Applications	5988-2164EN
Simultaneous stereoselective analysis of tramadol and its main phase I metabolites	
by online CZE electrospray ionization mass spectrometry	5980-1533E
A Comparison of CE-MS and LC-MS for Peptide Samples	5980-1304E
Implementing 21 CFR Part 11 in Analytical Laboratories Part 3: Ensuring Data Integrity in Electronic Records	5980-1305E
Implementing 21 CFR Part 11 in Analytical Laboratories Part 2: Security Aspects for Systems and Applications	5980-1306E
Implementing 21 CFR Part 11 in Analytical Laboratories Part 1: Overview and Requirements	5980-1308E
Capillary electrophoresis for the determination of forensic anions in adulterated foods and beverages	5968-9463E
Simultaneous determination of inorganic anions, organic acids, amino acids and carbohydrates by capillary electrophoresi	s 5968-4470E
Capillary preconditioning for analysis of anions using indirect UV detection in CZE.	
Systematic investigation of alkaline and acid prerinsing technquies by designed experiments	5968-3329E
Simultaneous Determination of Monosaccharides in Glycoproteins by Capillary Electrophoresis	5968-0772E
CE-ESI-MS of small molecules	5966-4661E
Instrumental Validation in Capillary Electrophoresis and Checkpoints for Method Validation	5965-9717E
Capillary electrophoretic determination of inorganic and organic anions using 2,6 pyridinedicarboxylic acid:	
effect of electrolyte's complexing ability	5965-8067E
Capillary Electrochromatography – a high efficiency micro-separation technique	5965-3430E
Micropreparative Single Run Fraction Collection of Peptides Separated by CZE for Protein Sequencing	5963-3506E

Technical Notes

Title	Publication No.
Capillary isoelectric focusing on the Agilent capillary electrophoresis system	5988-0974EN
CE-ESI-MS: An Integrated Solution	5980-2205E
Agilent ChemStation for GC, LC, LC/MSD, CE, UV-Visible and A/D Systems — Revision A.08.0x Enhancements	5980-1090E
Capillary Thermostatting in Capillary Electrophoresis	5968-8816E
HP ChemStation for GC, LC, LC/MSD, CE, UV-Visible and A/D Systems – Revision A.07.0x enhancements	5968-6668E
Use of Mobilities for Improved Reproducibility in CE	5968-2232E
High sensitivity Detection Cell for the Agilent Capillary Electrophoresis System	5965-5984E
Agilent Capillary Electrophoresis System – Technical Description	5965-6512E
Oligonucleotide Analysis – Optical filter for Polyacrylamide-filled Capillaries	5963-9870E
Improved Reproducibility & Increased Sample Throughput in Capillary Electrophoresis	5963-3296E
Use of Extremely Short Effective Length Capillaries in CE – Injection of Sample from the Outlet End of the Capillary	5963-3403E
Diode-Array Detection in Capillary Electrophoresis – Part 1: Using HP Extended Light Path Capillaries	5963-1889E
Diode-Array Detection in Capillary Electrophoresis – Part 2: Spectral Fidelity, Peak Purity, and Library Searching	5963-1891E

Agilent Technologies: Quality CE Support and Applications Literature

Articles in Scientific Journals

Automated peptide fraction collection in CE, Marzell Herold and Shiaw-lin Wu, LC-GC, Vol 12 Number 7, July 1994, pages 531-533

Purification and characterization of aspartate aminotransferase from E. coli by chromatographic methods. Marzell Herold, et al, International Journal of BioChromatography, 1994, Vol 1, pp29-45

Strategies for the development and validation of a CE method for enantiomeric purity testing of a basic drug.

Herbert Godel, poster presented at International Symposium for CE, York University, UK, 24-26 Aug 1994

Micropreparative single run fraction collection of peptides separated by CZE for protein sequencing.

Rudolf Grimm and Marzell Herold, Journal of Capillary Electrophoresis, July/August 1994

Identification of Tryptophan and Tyrosine residues in peptides separated by Capillary Electrophoresis by their second derivative spectra using diode-array detection. Rudolf Grimm, Arno Graf, David Heiger, ournal of Chromatography A, 679(1994)173-180

Micropreparative capillary isoelectric focusing of protein and peptide samples followed by protein sequencing. Rudolf Grimm, Journal of capillary electrophoresis, 002: 3 1995 (111-115)

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Basic Capillary Electrophoresis Troubleshooting

Symptom	Possible Cause	Solution(s)
Unstable Current		
Variable or no current	Air bubble formed in capillary	Flush capillary, ramp voltage to limit initial heating, and/or degas buffers.
	Clogged capillary	Flush capillary with absorbing solution (such as NaOH). A "step" in the base line should be observed when viewing the on-line signal at 200 nm. If still plugged, flush manually with syringe or high pressure gas.
	Broken capillary	Replace capillary.
	No or incorrect solution in buffer vials	Fill/change buffer vials.
	Large volume injection	Normal situation. Current should stabilize during analysis.
Unstable Baselin	10	
Spikes in baseline	Precipitates in buffer	Filter buffer through 0.2 or 0.45 µm filter.
	Micro air bubbles in buffer	Degas buffer by ultrasonication or vacuum.
	Precipitation of sample	Verify that sample components are sufficiently soluble in buffer.
Noisy baseline	Optical slit in capillary interface is occluded	Clean slit with methanol or water. View under magnifier.
	Aging deuterium lamp	Use DAD test to measure lamp ouput and time-on. Replace if necessary.
	Data acquisition rate too high	Determine peak width and decrease acquisition rate if appropriate.
	Improper reference wavelength	Acquire UV spectrum during analysis. Use lowest wavelength possible without impinging where sample absorbs. Also use wide bandwidth.
	Buffer absorbs at detection wavelength	Use minimally UV-absorbing buffers such as phosphate and borate, especially below 210 nm.
Drifting baseline	Improper capillary alignment	Re-seat capillary cartridge in detector block.
	Unequilibrated temperature	Allow 10-20 minutes for equilibration after opening top cover.
	Lamprecentlyignited	Allow 15-30 minutes for equilibration after igniting lamp.

Low Signal

Lowsignal	Sample concentration too low	Increase sample concentration.
	Insufficient sample stacking	Increase stacking by increasing difference between buffer and sample conductivities.
	Detection wavelength not optimized	Acquire UV spectrum during analysis. Use absorption maximum and appropriate bandwidth.
	Capillary id too narrow	Use Extended Light Path capillary

Poor Peak Efficiency

Broad peaks	Sample overloading	Decrease sample injection or concentration.
	Excessive Joule heating	Reduce voltage, buffer conductivity, or capillary id.

Symptom	Possible Cause	Solution(s)	
Poor Peak Efficiency			

Skewed peaks	Mismatched sample/buffer ion mobilities	Match mobilities or increase difference between buffer and sample conductivity.
	Sample overloading	Decrease sample injection or concentration.
Tailing peaks	Adsorption to capillary wall	Use pH extremes, high buffer concentrations, polymer additives, or coated capillary.

Poor Migration Time Reproducibility

3		
Adsorption to capillary walls	Changes in EOF caused by buffer (especially phosphates and detergents) or sample adsorption	Condition capillary and allow sufficient equilibration time. Replace capillary.
Hysteresis of wall charge	Caused by conditioning capillary at high (or low) pH and employing a low (or high) pH running buffer	Avoid pH differences. Allow sufficient equilibration time.
Changes in buffer	pH changes due to electrolysis	Replenish buffer.
composition	Buffer evaporation	Tightly cap buffer vials and reduce carousel temperature.
	Conditioning waste flushed into outlet reservoir	Use separate vial to collect waste.
	Conditioning solution carried over into buffer vial	First dip capillary in separate buffer or watervial.
Buffer reservoirs not level	Generation of laminar flow	Level liquid in reservoirs. If not replenishing buffer, do not use inlet vial for flushing capillary.
Different silanol content of capillary batches	Different wall charge and variations in EOF	Measure EOF and normalize.
Temperature changes	Changes in viscosity and EOF	Use system with capillary thermostating.

Poor Peak Area Reproducibility

Sudden application of high voltage	Heating, thermal expansion of buffer, and expulsion of sample	Ramp separation voltage or inject buffer plug after sample.
Sample evaporation	Increasing sample concentration and peak area	Cap vials and/or reduce temperature of sample carousel.
Instrumental limitations	System rise time significant proportion of injection time	Increase injection time.
Sample carry-over	Extraneous injection	Use capillary with flat, smooth injection end. Remove polyimide from end of capillary.
Zero-injection caused by simply dipping the capillary in the sample	Extraneous injection	Cannot be totally eliminated. Increase injection amount to minimize effect.
Sample adsorption to capillary walls	Distorted peak shape (tailing) Non-eluting sample	Change buffer pH. Increase buffer concentration. Use additive such as cellulose or coated capillary.
Low signal-to-noise ratio	Integration errors	Optimize integration parameters. Increase sample concentration. Use peak height.
Temperature changes of capillary environment	Changes in viscosity and injection amount	Use system with capillary thermostating.

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