Waters™

Note d'application

Pharmaceuticals and Personal Care Products in Water, Soil, Sediment and Biosolids by HPLC-MS/MS

Waters Corporation

Abstract

In this Application, we monitor the presence of pharmaceuticals in drinking water supplies and examining their long term effects on human health.

Introduction

Many hundreds of active compounds are used in both human and veterinary drug formulations. Due to the many different applications related to pharmaceuticals, their residues can reach the environment in multiple ways including excretion and manufacturing discharge. These compounds are not completely eliminated via sewage treatment plants, thus they can reach surface and groundwater supplies. Recently, there has been increased interest in monitoring for the presence of pharmaceuticals in drinking water supplies and examining their long term effects on human health.

Experimental

HPLC conditions

Instrument: Waters 2690 HPLC or Waters 2795 HPLC,

Quattro Ultima MS/MS

LC Column: Waters XTerra C_{18} , 3.5 μ m, 10.0 cm, 2.1 mm

Ionizatoin: Electrospray Positive (ES+)

Acquisition: MRM mode, unit resolution

Injection Volume: 15 μ L

LC G	radient Program	LC Flow Rate		General LC Cor	nditions
Time (min)	Flow Mixture ¹	(mL/min)	Gradient	Column Temperature	40 °C
0.0	95% Solvent A 5% Solvent B	0.150	1	Flow Rate	0.15 – 0.30 mL/min
4.0	95% Solvent A 5% Solvent B	0.250	6	Max Pressure	345 Bar
22.5	12% Solvent A 88% Solvent B	0.300	6	Autosampler Tray Temperature	4 °C
23.0	100% Solvent B	0.300	6	MS Conditi	ions
26.0	100% Solvent B	0.300	6	Source Temperature	140 °C
26.5	95% Solvent A 5% Solvent B	0.150	6	Desolvation Temperature	350 °C
33.0	95% Solvent A 5% Solvent B	0.150	6	Cone/Desolvation Gas Rate	80 L/hr /400 L/hr

¹ Solvent A = 0.3% Formic Acid and 0.1% Ammonium Formate in HPLC water

Solvent B = 1:1 Acetonitrile:Methanol

 $Group\ 1-Acidic\ extraction,\ positive\ electrospray\ ionization\ (ESI+)\ instrument\ conditions.$

					Detection	n Limits and	Minimum I	Levels	tract (ng/ΦL) Ation (ESI+) 2.2 12.5 0.9 1.25 5.7 50 0.1 1.25 0.4 2.5 0.1 0.5 0.2 2.5 0.1 0.5 0.2 1.25 0.2 1.25 0.1 0.5 0.3 1.25 0.1 0.5 0.3 1.25 0.1 0.5 1.25 0.1 0.5 0.2 1.25 0.1 0.5 0.1 0.5 0.2 1.25 0.1 0.5 0.3 1.25 0.1 0.5 0.1 0.5 0.1 0.5 0.1 0.5 0.2 1.25 0.3 1.25 0.1 0.5 0.3 1.25 0.4 1.25 0.1 0.5 0.3 1.25 0.4 1.25 0.5 1.7 5 1.7 5 1.8 5 1.7 5 1.9 0.2 0.5 1.9 0.1 0.5 1.9 0.2 0.5 1.9 0.1 0.5 1.9 0.
	RT	Parent-	Quantitation	Water	(ng/L)	Other (μg/kg)	Extract	(ng/ΦL)
Analyte	(min)	Daughter M/ZS	Reference	MDL	ML	MDL	ML	MDL	ML
Group 1	Ana	alytes Extracted l	Jnder Acidic Condition	ns and Ana	lyzed Using	Positive El	ectrospray l	onization ((ESI+)
Native Compounds						***	30000		
Sulfanilamide	2.5	190.0 - 155.8	13C ₆ -Sulfamethazine	8.9	50	48	200	2.2	12.5
Cotinine	2.8	177.0 - 98.0	Cotinine-d ₃	3.4	5	1.1	5	0.9	1.25
Acetaminophen	4.6	152.2 – 110.0	¹³ C ₂ - ¹⁵ N- Acetaminophen	27	200	35	200	6.7	50
Sulfadiazine	6.0	251.2 - 156.1	13C ₆ -Sulfamethazine	0.4	5	2.7	10	0.1	1.25
1,7-Dimethylxanthine	6.9	181.2 – 124.0	¹³ C ₃ -Caffeine	120	500	270	1000	30	125
Sulfathiazole	7.7	256.3 – 156.0	13C6-Sulfamethoxazole	0.5	5	1.9	50	0.1	1.25
Codeine	8.3	300.0 - 152.0	¹³ C ₃ -Trimethoprim	1.5	10	3.4	10	0.4	2.5
Sulfamerazine	8.7	265.0 - 156.0	¹³ C ₆ -Sulfamethazine	0.3	2	1.4	5	0.1	0.5
Lincomycin	9.3	407.5 - 126.0	¹³ C ₃ -Trimethoprim	0.8	10	4.7	10	0.2	2.5
Caffeine	9.3	195.0 – 138.0	¹³ C ₃ -Caffeine	15	50	5.4	50	3.6	12.5
Sulfamethizole	10.0	271.0 - 156.0	13C ₆ -Sulfamethoxazole	0.4	2	0.88	5	0.1	0.5
Trimethoprim	10.0	291.0 – 230.0	¹³ C ₃ -Trimethoprim	1.1	5	3.3	10	0.3	1.25
Thiabendazole	10.0	202.1 – 175.1	Thiabendazole-d ₆	0.7	5	2.1	10	0.2	1.25
Sulfamethazine	10.1	279.0 – 156.0	13C6-Sulfamethazine	0.6	2	0.83	5	0.2	0.5
Cefotaxime	10.2	456.4 - 396.1	¹³ C ₃ -Trimethoprim	10	20	18	50	2.5	5
Carbadox	10.5	263.2 - 231.2	¹³ C ₃ -Trimethoprim	2.3	5	2.1	10	0.6	1.25
Ormetoprim	10.5	275.3 – 259.1	¹³ C ₃ -Trimethoprim	0.3	2	0.50	2	0.1	0.5
Norfloxacin	10.7	320.0 - 302.0	¹³ C ₃ ¹⁵ N-Ciprofloxacin	28	50	15	50	7.0	12.5
Sulfachloropyridazine	10.8	285.0 - 156.0	13C ₆ -Sulfamethazine	1.2	5	1.9	5	0.3	1.25
Ofloxacin	10.8	362.2 - 318.0	13C ₃ 15N-Ciprofloxacin	1.8	5	3.4	10	0.4	1.25
Ciprofloxacin	10.9	332.2 - 314.2	13C ₃ 15N-Ciprofloxacin	5.1	20	8.1	20	1.3	5
Clinafloxacin	12.2	366.3 – 348.0	13C ₃ 15N-Ciprofloxacin	6.9	20	14	50	1.7	5
Digoxigenin	12.6	391.2 – 355.2	¹³ C ₃ -Trimethoprim	5.7	20	9.4	20	1.4	5
Oxolinic acid	13.1	261.8 - 243.8	¹³ C ₃ -Trimethoprim	0.6	2	0.62	2	0.2	0.5
Sulfadimethoxine	13.2	311.0 - 156.0	13C6-Sulfamethoxazole	0.1	1	0.55	2	0.03	0.25
Diphenhydramine	14.5	256.8 - 168.1	¹³ C ₃ -Trimethoprim	0.4	2	0.66	2	0.1	0.5
Penicillin G	14.6	367.5 - 160.2	¹³ C ₃ -Trimethoprim	2.4	10	13	50	0.6	2.5
Azithromycin	14.8	749.9 – 591.6	¹³ C ₃ -Trimethoprim	1.3	5	1.6	5	0.3	1.25
Flumegine	15.2	262.0 – 173.7	¹³ C ₃ -Trimethoprim	2.7	5	1.4	5	0.7	-
Ampicillin	15.3	350.3 – 160.2	¹³ C ₃ -Trimethoprim	- 1	5	-	5	-	1.25
Diltiazem	15.3	415.5 – 178.0	¹³ C ₃ -Trimethoprim	0.6	2	0.30	2	0.2	0.25
Carbamazepine	15.3	237.4 - 194.2	¹³ C ₃ -Trimethoprim	1.4	5	1.6	5	0.4	1.25
Penicillin V	15.4	383.4 – 160.2	¹³ C ₃ -Trimethoprim	4.4	20	19	50	1.1	5
Erythromycin	15.9	734.4 – 158.0	¹³ C ₂ -Erythromycin	-	1	-	2		0.25
Tylosin	16.3	916.0 – 772.0	13C ₂ -Erythromycin anhydrate	13	50	8.1	50	3.2	5
Oxacillin	16.4	434.3 – 160.1	¹³ C ₃ -Trimethoprim	3.3	10	9.4	20	0.8	2.5
Dehydronifedipine	16.5	345.5 – 284.1	¹³ C ₃ -Trimethoprim	0.6	2	0.41	2	0.2	0.5

Continuation of above table

					Detecti	ion Limits and	Minimum	Levels	evels			
	RT	Parent-	Quantitation	Water	(ng/L)	Other (μg/kg)	Extract	1.25			
Analyte	(min)	Daughter M/ZS	Reference	MDL	ML	MDL	ML	MDL				
Group 1	Ana	alytes Extracted	Under Acidic Condition	ons and An	alyzed Usir	ng Positive Ele	ectrospray	lonization ((ESI+)			
Native Compounds												
Clarithromycin	17.5	748.9 – 158.2	13C2-Erythromycin anhydrate	1.0	5	1.2	5	0.3	1.25			
Labeled compound	s spiked	into each sample	,	ė.	i.	Tree .						
Cotinine-d ₃	2.8	180.0 – 79.9	¹³ C ₃ Atrazine									
¹³ C ₂ - ¹⁵ N- Acetaminophen	4.5	155.2 – 111.0	¹³ C ₃ Atrazine									
¹³ C ₃ Caffeine	9.3	198.0 – 140.0	¹³ C ₃ Atrazine									
Thiabendazole-d ₆	9.8	208.1 – 180.1	¹³ C ₃ Atrazine									
¹³ C ₃ -Trimethoprim	10.0	294.0 – 233.0	¹³ C ₃ Atrazine									
¹³ C ₆ Sulfamethazine	10.1	285.1 – 162.0	¹³ C ₃ Atrazine									
¹³ C ₃ ¹⁵ N-Ciprofloxacin	10.9	336.1 – 318.0	13C3 Atrazine									
¹³ C ₆ -Sulfamethoxazole	11.2	260.0 – 162.0	13C ₃ Atrazine									
¹³ C ₂ -Erythromycin	15.9	736.4 – 160.0	13C ₃ Atrazine									
Fluoxetine-d ₅	16.8	315.3 – 153.0	13C ₃ Atrazine									
¹³ C ₂ -Erythromycin anhydrate	17.7	718.4 – 160.0	¹³ C ₃ Atrazine									
Injection internal s	tandard			10				-10				
¹³ C ₃ Atrazine	15.9	219.5 – 176.9 (134.0)	External standard									

Group 1 acidic extraction, positive electrospray ionization (ESI+) compound retention times (RTs), parent-daughter transitions, quantitation references, method detection limits, and minimum levels of quantitation.

HPLC conditions

Instrument:	Waters 2690 HPLC or Waters 2795 HPLC,
	Quattro Ultima MS/MS
LC Column:	Waters XTerra C ₁₈ , 3.5 μm, 10.0 cm, 2.1 mm
lonization:	Negative Ion Electrospray
ionization.	Negative for Electrospray
Acquisition:	MRM mode, unit resolution
Injection Volume:	5 μL

LC	Gradient Program	LC Flow Rate		General LC Con	ditions
Time (min)	Flow Mixture ¹	(mL/min)	Gradient	Column Temperature	40 °C
0.0	10% Solvent A 90% Solvent B	0.20	1	Flow Rate	0.20 – 0.23 mL/min
1.0	10% Solvent A 90% Solvent B	0.20	6	Max Pressure	345 Bar
18.0	40% Solvent A 60% Solvent B	0.23	6	Autosampler Tray Temperature	4 °C
20.0	90% Solvent A 10% Solvent B	0.23	6	MS Conditi	ons
24.0	90% Solvent A 10% Solvent B	0.23	6	Source Temperature	120 °C
24.3	10% Solvent A 90% Solvent B	0.20	6	Desolvation Temperature	400 °C
28	10% Solvent A 90% Solvent B	0.20	6	Cone / Desolvation Gas Rate	70 L/hr /450 L/hr

 $^{^{1}}$ Solvent A = 1:1 acetonitrile:methanol, with 5 mM Oxalic Acid

Solvent $B = HPLC H_2O$, with 5 mM Oxalic Acid.

Group 2 - Acidic extraction positive electrospray ionization (ESI+) instrument conditions.

					Detecti		ion Limits and Minimum Levels			
	RT	Parent-Daughter	Quantitation	Water (ng/L)		Other (ng/g)		Extract (ng/µL)		
Analyte	(min)	M/ZS	Reference	MDL	ML	MDL	ML	MDL	ML	
Group 2	Analy	tes Extracted Und	ler Acidic Condition	s and Ana	lyzed Usi	ng Positiv	e Electrosp	ray lonizati	on (ESI+)	
Native Compounds										
Minocycline	5.1	458.0 – 441.0	Thiabendazole-d ₆	51	200	-	200	13	50	
Epitetracycline	8.1	445.2 – 410.2	Thiabendazole-d ₆	3.6	20	8.6	20	0.9	5	
Epioxytetracycline (EOTC)	8.6	461.2 – 426.2	Thiabendazole-d ₆	4.1	20	18	50	1.0	5	
Oxytetracycline (OTC)	9.4	461.2 – 426.2	Thiabendazole-d ₆	2.1	20	2.2	20	0.5	5	
Tetracycline (TC)	9.9	445.2 – 410.2	Thiabendazole-d ₆	1.9	20	2.8	20	0.5	5	
Demeclocycline	11.7	465.0 – 430.0	Thiabendazole-d ₆	6.6	50	7.9	50	1.7	12.5	
Isochlortetracycline (ICTC) 1	11.9	479.0 – 462.2	Thiabendazole-d ₆	1.7	20	3.5	20	0.4	5	
Epichlortetracycline (ECTC) 1	12.0	479.0 – 444.0	Thiabendazole-d ₆	7.7	50	26	100	1.9	12.5	
Chlortetracycline (CTC)	14.1	479.0 – 444.0	Thiabendazole-d ₆	1.2	20	2.3	20	0.3	5	
Doxycycline	16.7	445.2 – 428.2	Thiabendazole-d ₆	2.8	20	2.3	20	0.7	5	
Epianhydrotetracycline (EATC)	17.0	426.8 – 409.8	Thiabendazole-d ₆	7.7	50	14	50	1.9	12.5	
Anhydrotetracycline (ATC)	18.8	426.8 – 409.8	Thiabendazole-d ₆	4.6	50	7.1	50	1.2	12.5	
Epianhydrochlortetracycline (EACTC)	20.7	461.2 – 444.0	Thiabendazole-d ₆	28	200	23	200	7.0	50	
Anhdrochlortetracycline (ACTC)	22.1	461.2 – 444.0	Thiabendazole-d ₆	5.2	50	11	50	1.3	12.5	
Labeled compound spiked	into ea	ch sample								
Thiabendazole-d ₆	7.0	208.1 – 180.1	13C ₃ Atrazine							
Injection internal standard	d					*				
¹³ C ₃ Atrazine	10.5	219.5 – 176.9 (134.0)	External standard							

 $^{^{1}}$ Isochlortetracycline (ICTC) is reported as the sum ICTC + ECTC due to a common transition ion.

Group 2 acidic extraction positive electrospray ionization (ESI+) compound retention times (RTs), parent-daughter transitions, quantitation references, method detection limits, and minimum levels of quantitation.

HPLC conditions

Instrument:	Waters 2690 HPLC or Waters 2795 HPLC,
	Quattro Ultima MS/MS
LC Column:	Waters XTerra C ₁₈ , 3.5 µm, 10.0 cm, 2.1 mm
Ionization:	Negative Ion Electrospray
Acquisition:	MRM mode, unit resolution
Acquisition	winivi illoue, utili resolution

LC	Gradient Program	LC Flow Rate		General LC Con	ditions
Time (min)	Flow Mixture ¹	(mL/min)	Gradient	Column Temperature	40 °C
0.0	60% Solvent A 40% Solvent B	0.2	1	Flow Rate	0.200 mL/min
0.5	60% Solvent A 40% Solvent B	0.2	6	Max Pressure	345 Bar
7.0	100% Solvent B	0.2	6	Autosampler Tray Temperature	4 C
12.5	100% Solvent B	0.2	6	MS Conditi	ons
12.7	60% Solvent A 40% Solvent B	0.2	6	Source Temperature	100 °C
16.0	60% Solvent A 40% Solvent B	0.2	1	Desolvation Temperature	350 ℃
				Cone/Desolvation Gas Rate	50L/hr /300 L/hr

 $^{^{1}}$ Solvent A = 0.1% Ammonium Acetate and 0.1% Acetic Acid in HPLC water

Solvent B = 1:1 MethanolAcetonitrile

Group 3 Acidic extraction negative electrospray ionization (ESI-) instrument conditions.

		Parent-			Detection	n Limits an	d Minimum	Levels	
	RT	Daughter	Ouantitation	Water	(ng/L)	Other ((Φg/g)	Extract	(ng/ΦL)
Analyte	(min)	M/ZS	Reference	MDL	ML	MDL	ML	MDL	ML
Group 3	Ana	lytes Extracted	Under Acidic Condition	s and Analy	zed Using I	Positive Ele	ctrospray l	onization ((ESI+)
Native Compounds									
Naproxen	6.7	228.9 – 168.6	13C-Naproxen-d ₃	3.9	10	6.1	20	1.0	2.5
Warfarin	7.1	307.0 - 117.0	Warfarin-d₅	0.9	5	1.6	5	0.2	1.25
Ibuprofen	8.4	205.1 – 161.1	¹³ C ₃ -lbuprofen	6.0	50	11	50	1.5	12.5
Gemfibrozil	9.5	249.0 - 121.0	Gemfibrozil-d₀	0.8	5	1.2	5	0.2	1.25
Triclocarban	9.6	312.9 – 159.7	13C ₆ -Triclocarban	2.1	10	2.7	10	0.5	2.5
Triclosan	9.7	286.8 – 35.0	13C ₁₂ -Triclosan	92	200	56	200	23	50
Labeled compounds	s spiked i	nto samples							
¹³ C-Naproxen-d ₃	6.6	232.9 – 168.6	¹³ C ₆ -TCPAA						
Warfarin-d₅	7.0	312.0 – 161.0	¹³ C ₆ -TCPAA						
¹³ C ₃ -lbuprofen	8.5	208.2 – 163.1	13C ₆ -TCPAA						
Gemfibrozil-d ₆	9.5	255.0 – 121.0	13C ₆ -TCPAA						
¹³ C ₆ -Triclocarban	9.6	318.9 – 159.7	13C ₆ -TCPAA						
¹³ C ₁₂ -Triclosan	9.7	298.8 – 35.0	¹³ C ₆ -TCPAA						
Injection Internal Star	ndard								
¹³ C ₆ -TCPAA	4.9	258.8 – 200.7	External standard						

Group 3 acidic extraction negative electrospray ionization (ESI-) compound retention times (RTs), parent-daughter transitions, quantitation references, method detection limits, and minimum levels of quantitation.

HPLC conditions

Instrument:	Waters 2690 HPLC or Waters 2795 HPLC, Quattro Ultima MS/MS
LC Column:	Waters Atlantis HILIC, 3.0 µm, 2.1x 100 mm
lonization:	Electrospray Positive (ES+)
Acquisition:	MRM mode, unit resolution
Purge Solvent:	100% CH ₃ CN (changed from H2O)
Injection Volume:	2.0 µL

LC	Gradient Program	LC Flow Rate		General LC Con	ditions
Time (min)	Flow Mixture ¹	(mL/min)	Gradient	Column Temperature	40 °C
0.0	2% Solvent A 8% Solvent B	0.25	1	Flow Rate	0.25 mL/min
5.0	30% Solvent A 70% Solvent B	0.25	6	Max Pressure	345 Bar
12.0	30% Solvent A 70% Solvent B	0.25	6	Autosampler Tray Temperature	4 °C
12.5	2% Solvent A 98% Solvent B	0.25	6	MS Conditi	ons
16.0	2% Solvent A 98% Solvent B	0.25	6	Source Temperature	120 °C
				Desolvation Temperature	350 °C
				Cone/Desolvation Gas Rate	70L/hr /400 L/hr

¹ Solvent A = 0.1% Acetic Acid/Ammonium Acetate Buffer

Solvent B = Acetonitrile

Group 4 - Basic extraction positive electrospray ionization (ESI+) instrument conditions.

		Parent-			Detecti	on Limits and	d Minimum	Levels		
	RT	Daughter M/	/ Quantitation W	Water	Water (ng/L)		Other (ng/g)		Extract (ng/ΦL)	
Analyte (min)	ZS	Reference	MDL	ML	MDL	ML	MDL	ML		
Group 4	Analı	tes Extracted U	Jnder Acidic Conditi	ons and Ana	lyzed Usin	g Positive El	ectrospray	lonization (+) ESI	
Native Compounds										
Cimetidine	6.9	253.1 – 159.0	Albuterol-d ₃	0.6	2	0.78	2	0.2	0.5	
Albuterol	9.4	240.0 – 148.0	Albuterol-d ₃	0.9	2	0.39	2	0.2	0.5	
Ranitidine	10.3	315.0 – 175.9	Albuterol-d ₃	0.7	2	1.1	2	0.2	0.5	
Metformin	11.0	131.1 – 60.1	Metformin-d ₆	23	100	38	100	5.8	25	
Labeled compounds	s spiked in	ito samples								
Albuterol-d ₃	9.4	243.0 – 151.0	Cotinine-d ₃							
Metformin-d ₆	11.0	285.1 – 163.0	Cotinine-d ₃							
Injection Internal S	tandard									
Cotinine-d ₃	5.9	180.0 – 79.9	External standard						\$\text{\$\exitt{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\exitt{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\exitt{\$\text{\$\exitt{\$\text{\$\}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}	
¹³ C ₃ -Atrazine	2.0	219.5 – 176.9 (134.0)	External Standard							

Group 4 basic extraction positive electrospray ionization (ESI+) compound retention times (RTs), parent-daughter transitions, quantitation references, method detection limits, and minimum levels of quantitation.

References

- 1. Total Solutions for Environmental Applications: 720002163EN
- 2. LC-MS Determination of Pharmaceutical Residues in Environmental Samples: 720000421EN

Featured Products

Alliance HPLC System https://www.waters.com/534293

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