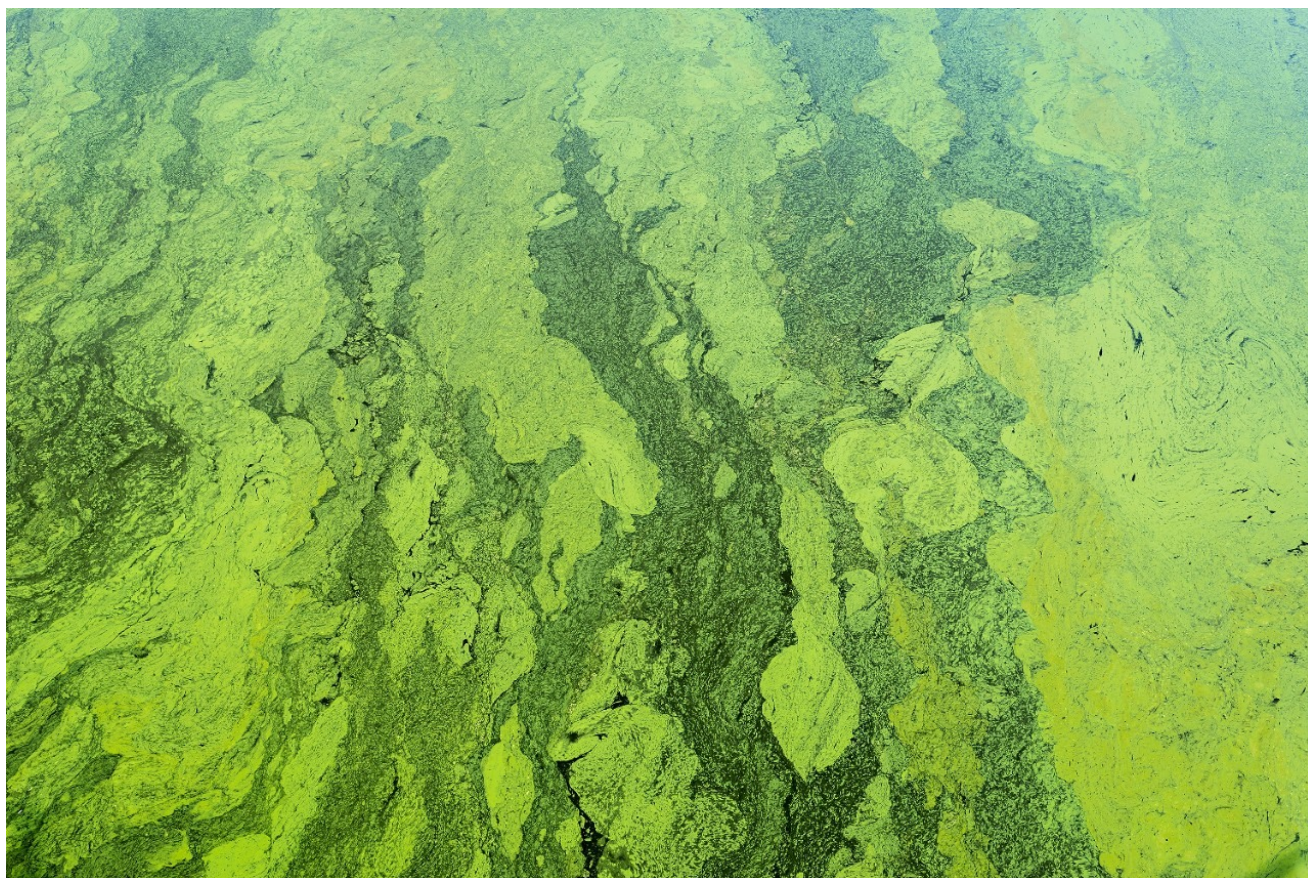


## Analysis of Microcystins in Urine with 2D-LC-MS/MS – Part III

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### Abstract

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This application demonstrates the effectiveness of captive SPE extraction process for microcystins by 2D-LC-

MS/MS from urine.

## Benefits

- Extraction protocol optimization <30 minutes.
- Effective elimination of time-consuming evaporation and reconstitution steps with 2D-LC-MS/MS.

## Introduction

Algae “super blooms” are a commonly encountered environmental issue in fresh water that occurs due to the buildup of cyanobacteria. Their presence is predominantly linked to excess nutrients (fertilizers) from water runoff. Many of the commonly encountered cyanobacteria produce potent cyanotoxins that pose serious health threats and even death to local wild life and humans. Microcystins are produced by the most notable and widespread cyanobacteria<sup>1,2</sup> “*Microcystis aeruginosa*” from which the toxins take their name. When released, microcystins may persist for weeks to months, because they are nonvolatile, hydrophilic, stable in sunlight, and stable over a wide temperature/pH range. Chemically, microcystins are cyclic heptapeptides produced through non-ribosomal peptide synthases. Microcystins consist of a seven-membered peptide ring which is made up of five non-protein amino acids and two protein amino acids. It is these two protein amino acids that distinguish microcystins from one another, while the other amino acids are more or less constant between variant microcystins. Using amino acid single letter code nomenclature, each microcystin is designated a name depending on the variable amino acids which complete their structure. Microcystin-LR is named for containing the variable amino acids leucine (L) and arginine (R). Microcystin-LR was the first identified and is the most commonly studied to date. Other common microcystins include RR, YR for Tyrosine (Y), and LA for Alanine (A).

Microcystins in drinking water are not regulated by the US EPA, however, they are unregulated microbial drinking water contaminants listed on the US EPA Contaminant Candidate Lists (CCLs) 1 and 2 as cyanobacteria and their toxins. Certain cyanotoxins, specifically anatoxin-A, microcystin-LR and cylindrospermopsin, are on the US EPA CCL3. The World Health Organization<sup>3</sup> (WHO) conducted an evaluation of the tolerable daily intake (TDI) level, based on a non-cancer endpoint. The obtained value of 0.04 micrograms per kilogram body weight per day ( $\mu\text{g}/\text{kg}/\text{d}$ ), is based on the results of liver toxicity studies in mice. A TDI is the maximum daily dose of microcystins that is considered safe. Using this TDI, the WHO also developed a drinking water concentration limit of 1.5  $\mu\text{g}/\text{L}$  for microcystin LR. For this study, they assumed that a 60 kg (132 lbs.) person drinks two liters of water each day and that 80% of the two liters is from a contaminated source. Furthermore, microcystin

contaminated fresh-water that empties into the ocean has been shown to also lethally affect marine life in the area of contamination.

Human consumption of tainted sea life can lead to microcystin exposure and even death. Several analytical techniques that have been developed for the analysis of microcystins include the mouse bioassay,<sup>4</sup> phosphatase inhibition<sup>5</sup> assay, enzyme-linked immunosorbent assay (ELISA),<sup>6</sup> and reversed-phase high performance liquid chromatography (RP-HPLC),<sup>7-8</sup> just to name a few. Little has been done with multi-dimensional chromatography, thus a method was developed for forensic postmortem analysis in biological fluids of microcystins RR, LR, and YR by 2-dimensional (2D) liquid chromatography mass spectrometry (LC-MS/MS).<sup>9-11</sup>

## Experimental

Two MRM transitions (quantification and confirmation) for each microcystin were selected and optimized. The MRM conditions are listed in Table 1. For this application, finding the optimum extraction and chromatographic conditions for this multi-residue analysis posed a significant challenge. The chromatographic conditions were tested on several trapping chemistries (Oasis HLB, XBridge C<sub>18</sub>, and XBridge C<sub>8</sub>) and one separation chemistry (BEH C<sub>18</sub>) from Waters Corporation, Milford, MA, USA. The loading (low pH, high pH, and neutral pH) and eluting mobile phase (methanol or acetonitrile at low and high pH) were also optimized using an automated 6 × 6 process. All microcystin standards were purchased from Enzo Lifesciences (Figure 1). All solvents were LC grade or higher and obtained from Fisher Scientific (Fair Lawn, NJ, USA). The formic acid and ammonium hydroxide were purchased from Sigma Aldrich (St. Louis, MO, USA). The extraction process was performed on pre-conditioned reversed-phase sorbent (Oasis HLB SPE 3 cc 60 mg barrel, Waters Corporation, Milford, MA, USA) for the captive extraction for the screening extraction.

Toxin	MW (g/mol)	Precursor	Cone	Quant	CID	Qual	CID
Microcystin LR	955.19	995.9	60	135.1	65	86.1	55
Microcystin RR	1038.2	520.0	30	135.1	25	70.1	50
Microcystin YR	1054.2	1045.9	60	135.1	60	70.1	80
Nodularin (IS)	824.98	825.7	60	135.1	50	70.1	65

Table 1. MRM transitions for YR, LR, and RR.

The image shows a complex chemical structure of a molecule, likely a peptide or a related compound. A red circle highlights a specific functional group: a p-hydroxybenzyl group, which consists of a benzene ring with a hydroxyl group (-OH) at the para position, attached to a methylene group (-CH<sub>2</sub>-). This group is connected to a carbonyl group (-C(=O)-) which is part of a larger chain. The rest of the molecule includes various amino acid residues, a guanidino group, and a methoxy group (-OCH<sub>3</sub>).

[illegible][illegible]

## Loading conditions

Oasis HLB 20  $\mu\text{m}$ –80 mg,  $2.1 \times 30$  mm

MilliQ water (pH 7, no additives)

2 mL/min

5% (0.1 mL/min loading pump and 2 mL/min diluting pump)

ACQUITY UPLC with 2D Technology configured  
for trap and elute with at-column dilution

10 min

ACQUITY UPLC BEH C<sub>18</sub>, 1.7 μm, 2.1 × 50 mm  
(p/n: 186002350)

Column temp.:	60 °C
Mobile phase A:	Water + 0.5% formic acid
Mobile phase B:	Acetonitrile + 0.5% formic acid
Elution:	5 min linear gradient from 5% (B) to 95% (B)
Flow rate:	0.500 mL/min (elution pump)
Injection volume:	100 µL

## MS conditions

System:	Xevo TQ-S
Ionization mode:	ESI+
Capillary voltage:	3.0 kV
Cone voltage:	90.0 V
Source temp.:	150 °C
Desolvation temp.:	550 °C
Desolvation gas:	1100 L/hr
Cone gas:	50 L/hr

## Results and Discussion

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### Separation and Detection



The separation conditions were optimized for each microcystin according to a 2D-LC 6 × 6 elution scheme (Figure 2) for both aqueous and organic standards. Microcystins are heptapeptides, and these molecule types have the potential to show zwitterionic behaviour and possibly lead to adsorption or solubility issues. When utilizing multidimensional chromatography, a wide range of options become available, not seen with single dimension separations. At-column dilution (ACD) allows for the option of injecting 100% organic extracts which in turn yields a 50% reduction during sample preparation. In Figure 3, a side-by-side comparison of a 15 µL acetonitrile injection in 1D versus 100 µL acetonitrile injection in 2D clearly shows the versatility of the multidimensional chromatography technique. The breakthrough effect of an organic extract in 1D is a known fact and has been well documented for years. Therefore, the large volume injection of a 100% organic extract capability of a 2D system with a trap-and-elute ACD configuration, can have a significant positive impact on sample preparation protocol.

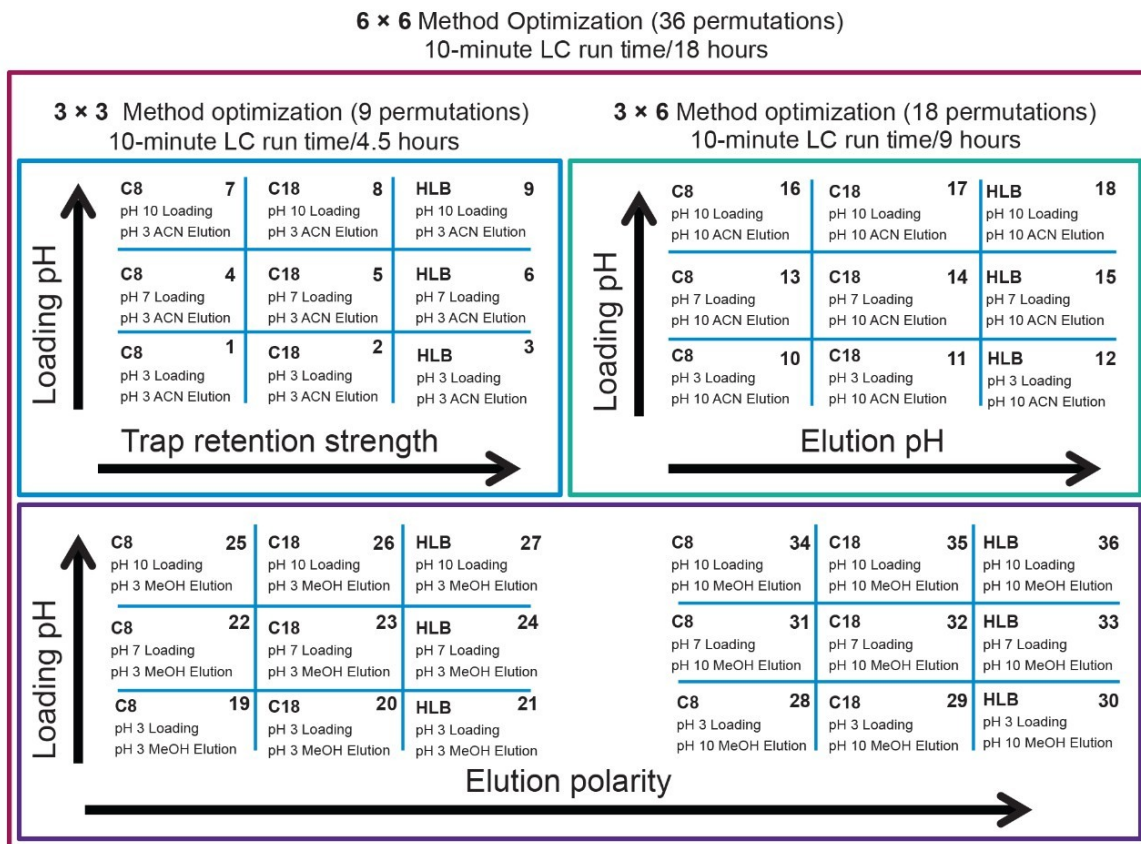


Figure 2. Grid for 6 x 6 method optimization.

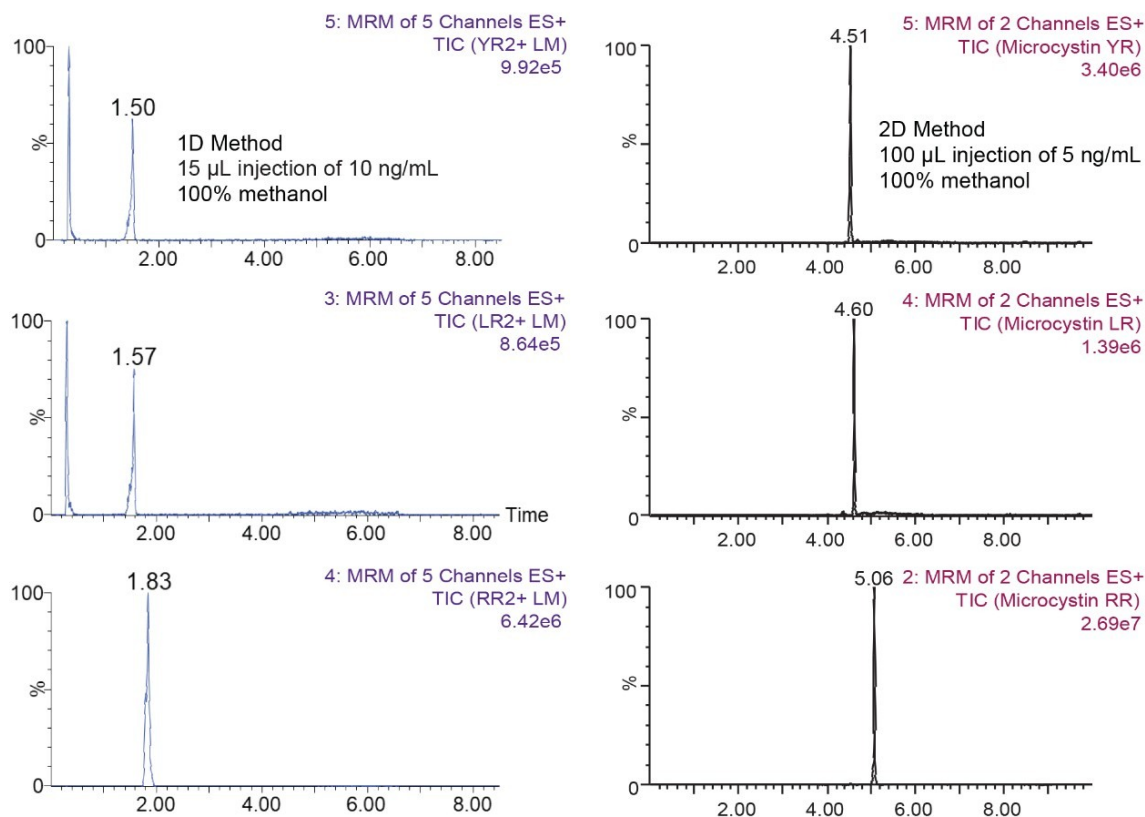


Figure 3. 1D vs 2D for YR, LR, and RR.

The results of the 6 × 6 grid are shown in Figure 4. With the optimization of four key parameters, a target analyte can be focused in a tight narrow band on the first dimension and transfer onto the second dimension with a Gaussian profile. The compilation of the 36-method variants gave an in-depth understanding of a target analyte's separation behaviour. The grid is colour coded to give a strong visualization effect to identify which condition will produce a Gaussian profile (green box). If a set of conditions produces a signal with any type of chromatography distortion (i.e., leading, split, tailing, shoulder, etc.), the corresponding result is flagged by a yellow box. All absence of signal is highlighted by a red box. Overall, microcystin YR shows that up to 25 methods (aqueous and organic standards) or 29 (organic standards only), out of 36 total, will yield a Gaussian profile. The rationale behind an aqueous versus methanol or acetonitrile standard is to investigate the possibility of adsorption or non-specific binding onto the glass surface, if standards are stored in glass vials and under aqueous conditions. Once all 36-method results are compiled for each microcystin, the next phase is to identify which method can be used for all three target analytes. In this instance, since there are only three analytes plus an internal standard, the results in Figure 5 cross reference all 36 methods against the methanol and acetonitrile standards for each microcystin. The compilation eliminated all yellow and red results and concentrated only on Gaussian profile (green box) and highest intensities. Each method is given a score, and for each method producing a maximum score, the result is highlighted in the far end column, listing which methods can be used for the application. In

this instance, 28 methods can be selected for the analysis of microcystins YR, LR, and RR, including nodularin as internal standard. For this application, method 14 was selected for highest signal intensities across all four analytes for both methanol and acetonitrile standards.

	1	2	3	4	5	6	7	8	9
Microcystin YR (H <sub>2</sub> O)	e5	e5	e5	blob	mp e4	blob	e6	e6	tail
Microcystin YR (MeOH)	e6	e6	e6	e6	e6	e6	e6	e6	tail
Microcystin YR (ACN)	e6	e6	e6	e5	e6	e5	e5	e6	tail
	10	11	12	13	14	15	16	17	18
Microcystin YR (H <sub>2</sub> O)	e5	e6	e5	e4	e5	e5	e6	e5	e5
Microcystin YR (MeOH)	e6	e7	e6	e6	e7	e6	e7	e6	e6
Microcystin YR (ACN)	e6	e6	e6	e6	e7	e6	e6	e6	e6
	19	20	21	22	23	24	25	26	27
Microcystin YR (H <sub>2</sub> O)	e4	mp e4	tail	mp e4	mp e4	mp e4	e6	e6	tail
Microcystin YR (MeOH)	e5	e6	tail	mp e4	e6	blob	e6	e6	blob
Microcystin YR (ACN)	e5	e5	tail	mp e4	e6	blob	e6	e6	blob
	28	29	30	31	32	33	34	35	36
Microcystin YR (H <sub>2</sub> O)	e5	e5	tail	e5	e4	tail	e6	e6	tail
Microcystin YR (MeOH)	e6	e6	tail	e7	e6	tail	e6	e6	tail
Microcystin YR (ACN)	e6	e6	tail	e6	e5	tail	e6	e5	tail

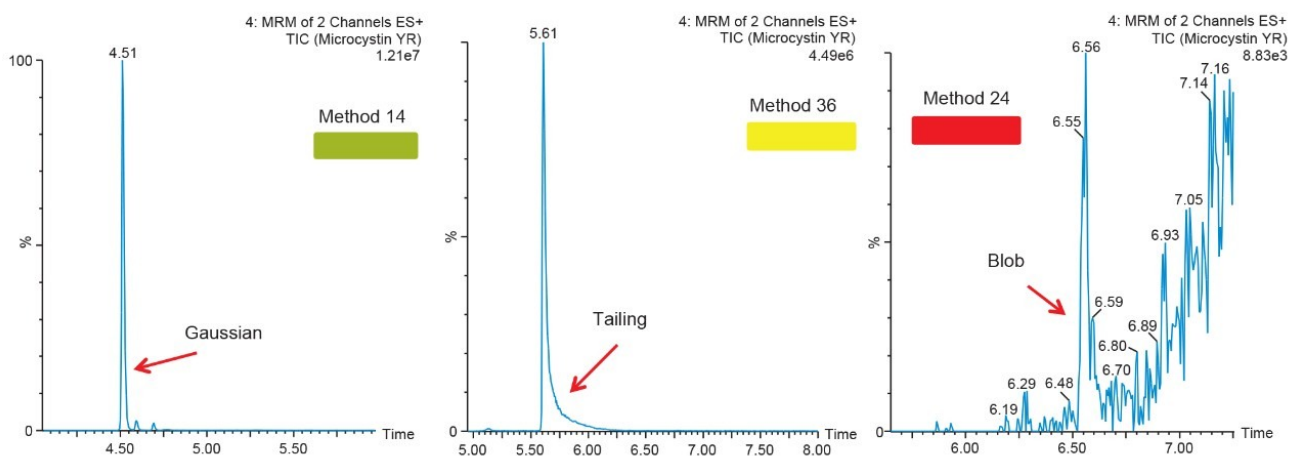


Figure 4. The 6 × 6 grid target hit for YR.



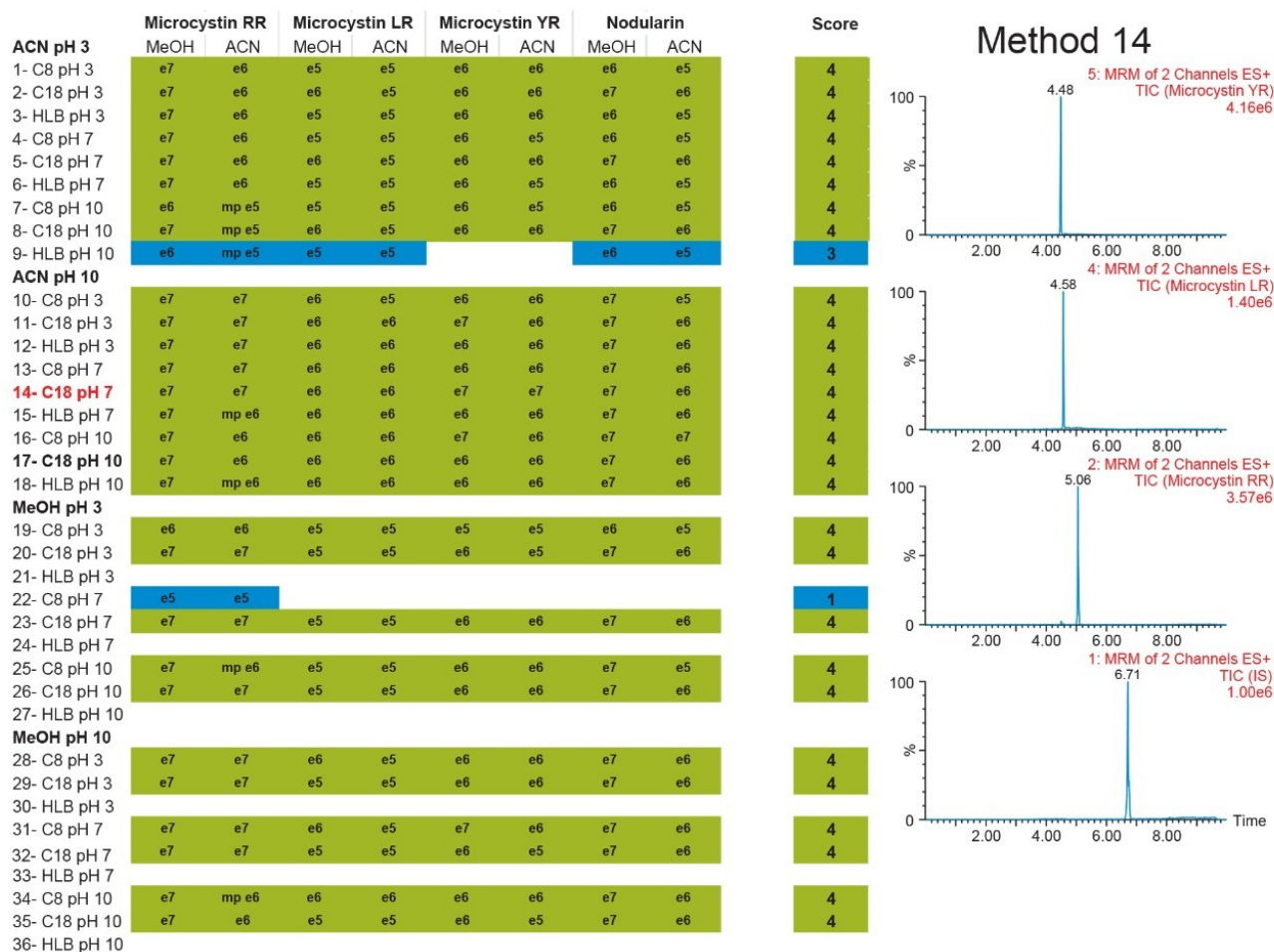


Figure 5. Cumulative 6 × 6 hit for YR, LR, and RR.

## Single Stage 2D Captive Extraction

Once the LC and MS optimization phases were completed, the next steps focused on sample cleanup, enrichment, and retention behavior. In this instance, mapping the retention behavior at the sample preparation level produces cleaner extracts. This strategy relies on identifying key elution points (0–100% recovery) against the target analyte's pKa and solubility. Figure 6 shows a threestep sequential elution, with three elution strengths and two pH values. The experimental workflow utilized two SPE cartridges for three elution solvents (methanol, acetonitrile, and acetone) with a 10% elution incremental from 0–100% at pH 3 and 10. Each elution cut was analyzed by the 2D-LC trap and elute with an ACD system without any need for solvent evaporation and reconstitution. The results are tabulated in Table 2. The rationale of using pH 3 and 10 was to understand the retention behavior of microcystins according to their pKa values. For example, under neutral elution conditions, it is expected to see high  $k'$  values, thus requiring a higher percentage of organic for a total release of the target analyte from the extraction sorbent and vice versa under ionized conditions. This separation behavior would fit if the target analyte is comprised of basic moieties. For microcystin RR, at pH 3 elution with acetonitrile, the

analyte is completely eluted between 20%–40%, with the bulk of the analyte eluting at 30%. However, at pH 10, RR elutes in a tighter elution cut between 20%–30%. This result suggests that RR behaves as an acidic entity (low  $k'$  at pH 3 and high  $k'$  at pH 10). The microcystins LR and YR are producing similar profiles under low and high pH elution, but at different elution profiles. The acetone and methanol showcase pH profiles but at higher elution percentages, thus giving insight on the microcystin's solubility.

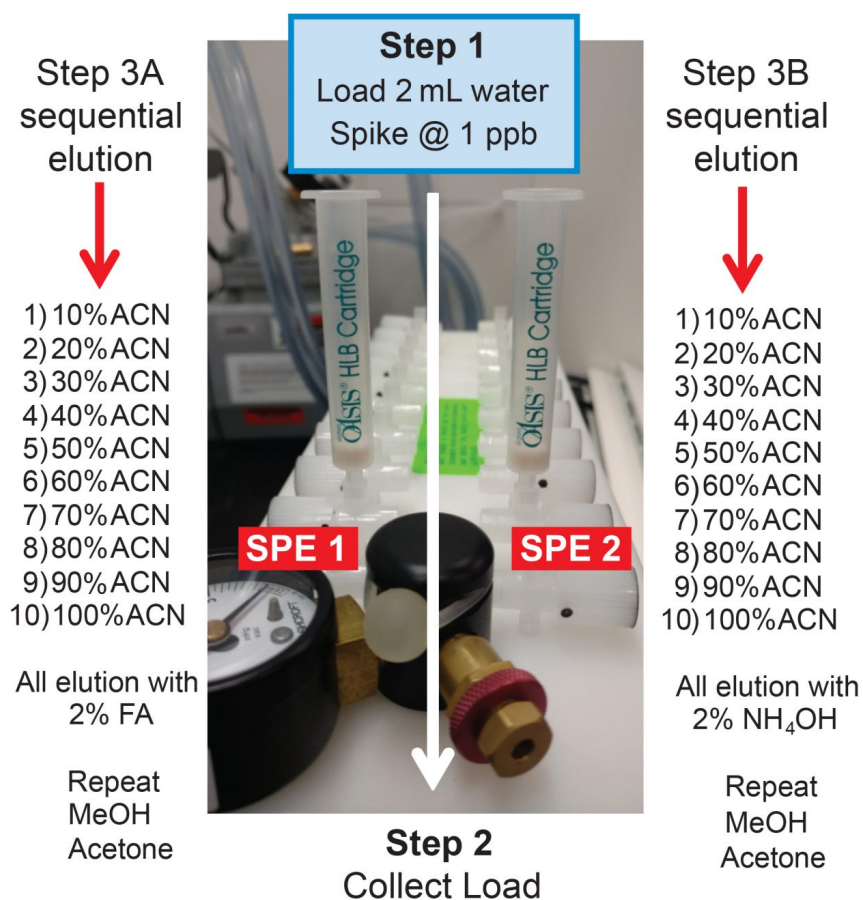


Figure 6. Single-stage 2D extraction protocol.



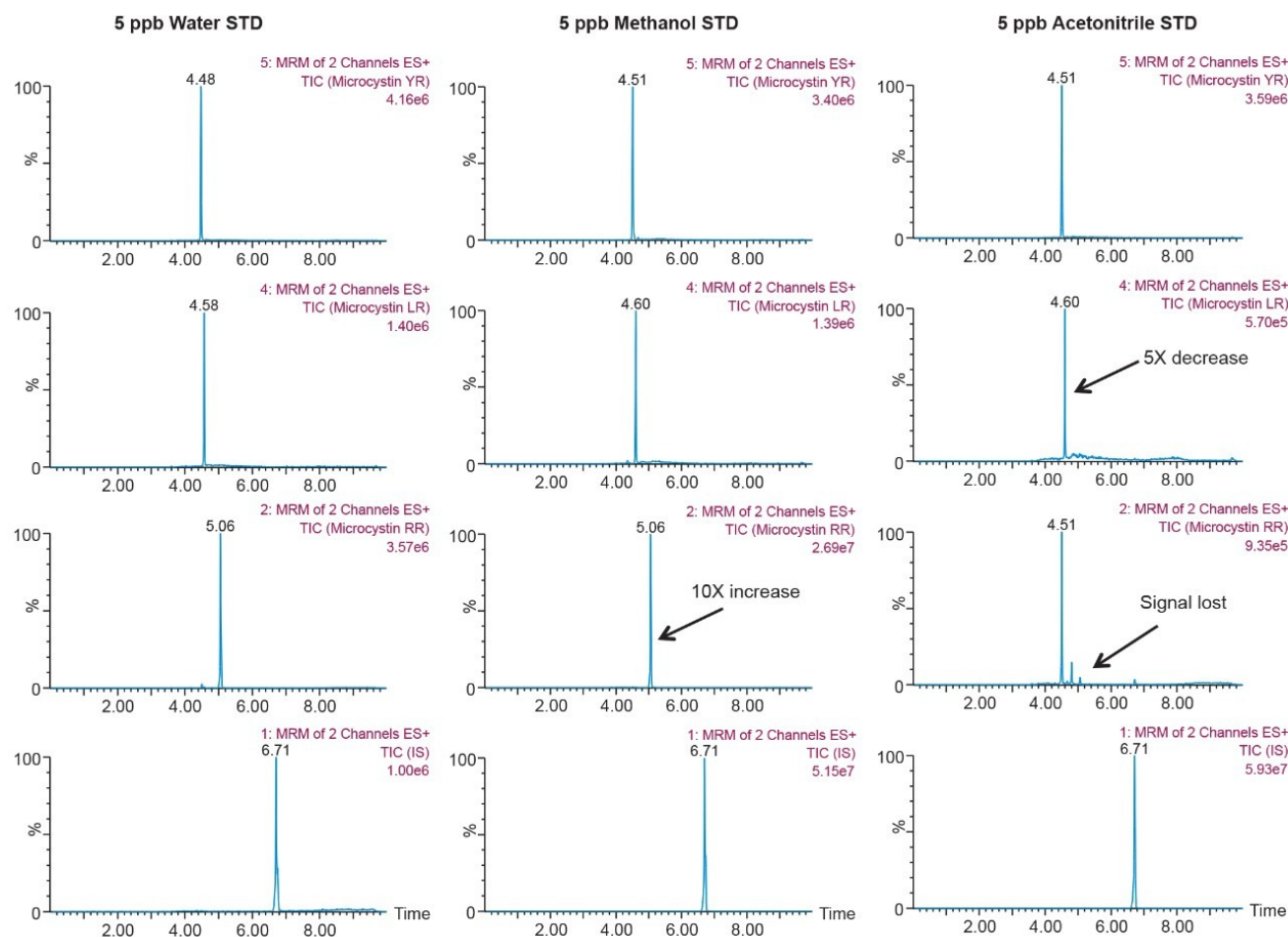


Figure 7. A 2D-LC-MS/MS chromatogram for YR, LR, and RR in water, methanol, and acetonitrile.

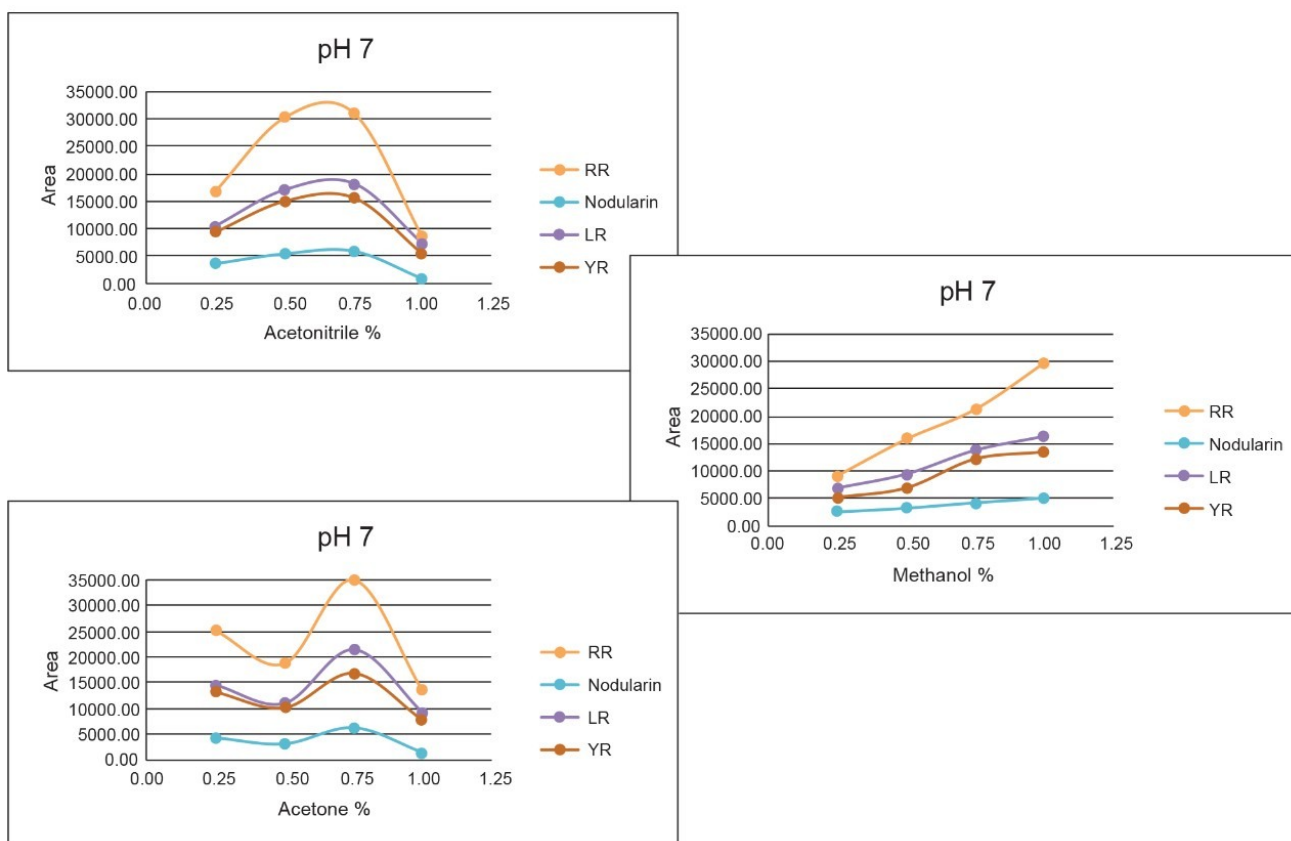


Figure 8. Elution profile of YR, LR, and RR with methanol, acetonitrile, and acetone.



			0%	10%	20%	30%	40%	50%	60%	70%	80%	90%	100%
RR	ACN	pH 3	0.11	0.23	16.86	64.69	14.65	1.59	0.83	0.45	0.31	0.20	0.08
		pH 7	0.07	0.12	0.13	0.16	0.28	21.14	61.17	14.39	1.61	0.56	0.38
		pH 10	0.09	0.37	15.17	76.88	5.72	0.64	0.39	0.28	0.21	0.23	0.02
	MeOH	pH 3	7.62	0.70	0.35	0.24	1.76	24.23	52.42	10.03	1.50	0.72	0.43
		pH 7	0.20	0.21	0.23	0.15	0.18	15.83	69.98	11.65	0.83	0.34	0.39
		pH 10	0.08	0.09	0.07	0.07	0.14	11.22	64.99	20.12	2.03	0.67	0.53
	Acetone	pH 3	0.05	0.21	5.39	66.69	18.85	3.32	1.59	1.20	1.04	0.92	0.73
		pH 7	0.02	0.23	11.74	70.21	13.22	2.37	0.91	0.25	0.26	0.41	0.37
		pH 10	0.10	3.28	6.28	76.83	8.42	3.47	0.67	0.32	0.12	0.44	0.07
LR	ACN	pH 3	0.28	0.24	0.44	59.39	34.14	3.57	0.63	0.30	0.34	0.33	0.34
		pH 7	0.35	0.85	4.53	13.25	8.05	15.34	32.21	21.22	2.99	0.65	0.55
		pH 10	0.45	1.21	88.17	7.97	0.85	0.50	0.15	0.28	0.21	0.19	0.02
	MeOH	pH 3	5.01	0.32	0.24	0.20	0.29	0.41	44.55	40.80	6.41	1.11	0.65
		pH 7	0.04	0.96	10.80	8.72	6.63	9.91	44.31	16.35	1.38	0.38	0.52
		pH 10	0.05	0.12	0.06	0.39	19.43	61.19	17.24	1.03	0.20	0.10	0.19
	Acetone	pH 3	0.15	0.09	0.07	5.05	79.15	14.40	0.10	0.19	0.21	0.34	0.23
		pH 7	0.06	51.04	32.64	11.79	2.97	1.02	0.27	0.15	0.02	0.03	0.03
		pH 10	0.27	15.16	57.77	25.54	1.06	0.10	0.01	0.01	0.01	0.04	0.02
YR	ACN	pH 3	0.28	0.32	0.44	50.39	41.14	5.07	0.92	0.45	0.36	0.31	0.34
		pH 7	0.33	0.42	2.91	11.04	7.75	14.95	31.56	25.17	4.39	0.91	0.58
		pH 10	0.60	7.07	84.28	5.99	0.71	0.57	0.20	0.15	0.15	0.19	0.09
	MeOH	pH 3	2.47	0.23	0.24	0.17	0.32	0.34	39.32	46.39	8.61	1.22	0.69
		pH 7	7.62	0.70	0.35	0.24	1.76	24.23	52.42	10.03	1.50	0.72	0.43
		pH 10	0.11	0.11	0.16	3.26	33.90	49.30	11.85	0.77	0.23	0.13	0.17
	Acetone	pH 3	0.06	0.06	0.08	3.69	74.40	18.92	1.66	0.51	0.19	0.22	0.19
		pH 7	0.01	46.13	28.25	17.94	5.41	1.71	0.45	0.02	0.04	0.02	0.03
		pH 10	0.182	31.286	60.929	7.119	0.365	0.021	0.013	0.012	0.021	0.041	0.012

Final conditions: Wash 1: 30% MeOH pH 3  
Wash 2: 30% MeOH pH 10  
Elution: 70% MeOH pH 10

Table 3. Elution profile of YR, LR, and RR with methanol, acetonitrile, and acetone.

The results for the first extraction method with a wash step at 30% methanol at pH 3 followed by an elution at 40% acetonitrile at pH 10 (Figure 9), showing drastic low recovery values for RR and LR when extracting from urine. This confirms the trend seen in Figure 8, therefore an updated method was chosen in which the elution solvent is methanol to avoid solubility issues during extraction. The final SPE conditions chosen comprised of two wash steps at 40% methanol at both pH 3 and pH 8, and an elution step of 70% methanol at pH 10, yielded excellent recoveries for YR, RR, and LR in urine (Figure 10).

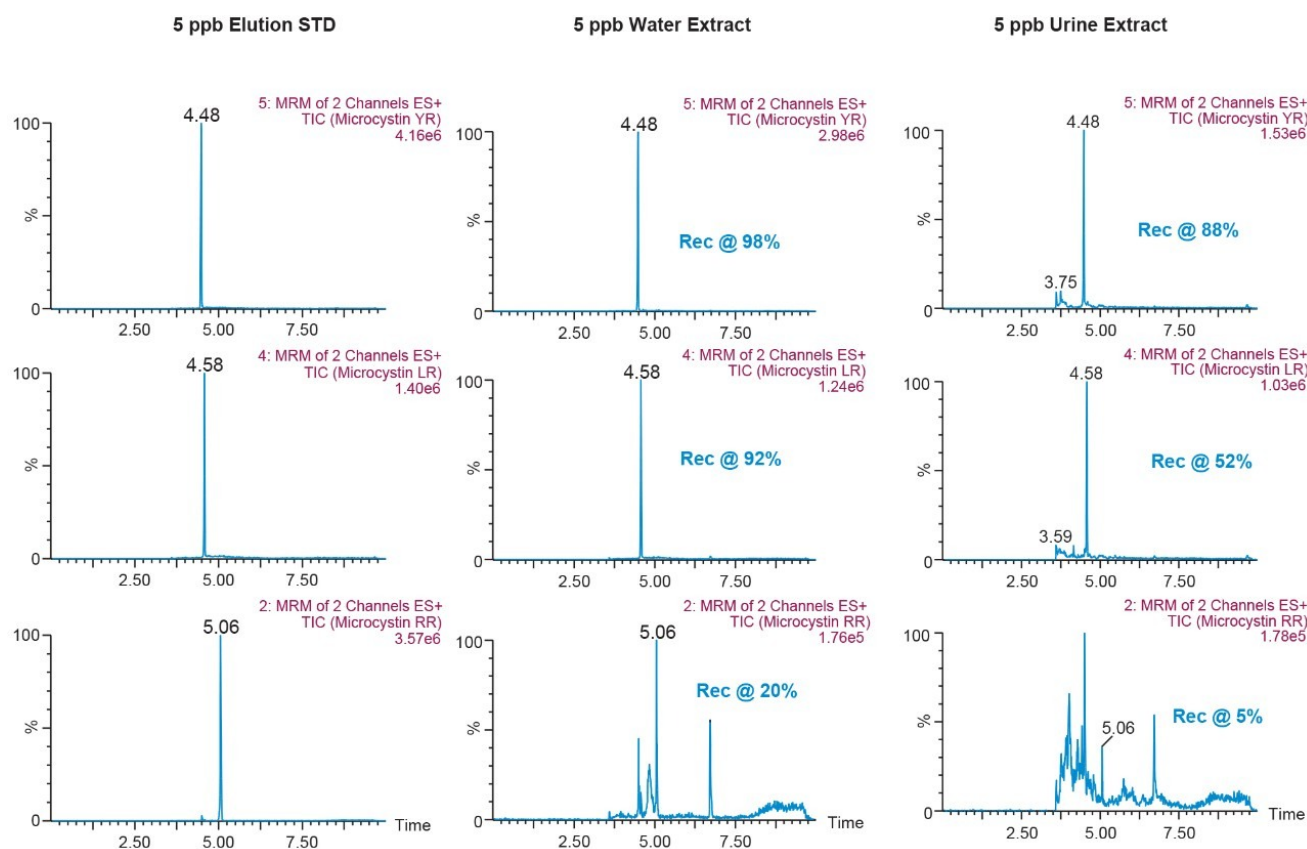


Figure 9. Extraction protocol 1 – wash at 30% methanol at pH 3 and elution with 40% acetonitrile at pH 10.

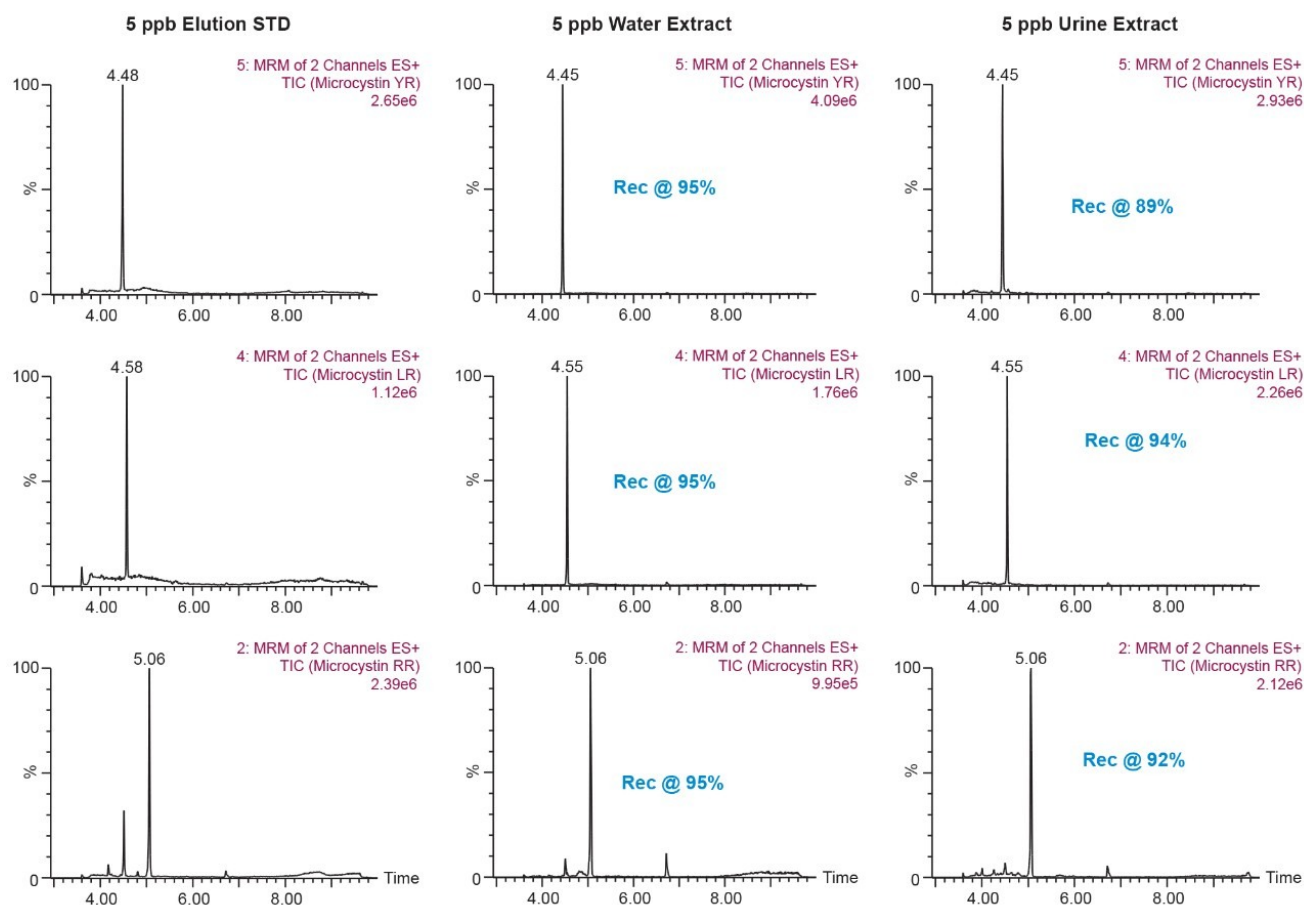


Figure 10. Extraction protocol 2 – wash with 40% methanol at pH 3 and pH 10; elution with 70% methanol at pH 10.

## Conclusion

This application demonstrates the effectiveness of captive SPE extraction process for microcystins by 2D-LC-MS/MS from urine. The  $6 \times 6$  chromatography method optimization was completed over ~18 hours. The option of direct large volume injections of up to 100% organic extracts made it possible to optimize an extraction protocol with a total work time of <30 minutes. This added benefit of 2D-LC-MS/MS allows direct injections of aqueous and organic extracts at various percentages to effectively eliminate time-consuming evaporation and reconstitution steps. The captive extraction protocol produced a 90% recovery of microcystins YR, LR, and RR in urine.

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