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# Development and application of an LC-MS method to the determination of poly- and perfluoroalkyl substances (PFASs) in drinking, sea and surface water samples†

Belinda Huerta, <sup>a</sup> Brendan McHugh<sup>b</sup> and Fiona Regan <sup>a</sup>

Poly- and perfluoroalkyl substances (PFASs) are a group of synthetic organic surfactants that have become a global concern because of their toxicity and widespread presence in the aquatic environment and organisms globally. In this study, a new analytical method has been developed and validated for the analysis of 15 perfluorinated compounds in different water matrices: river water, drinking water and seawater. Water extraction was performed in anion exchange solid phase extraction cartridges, and extracts were analysed by liquid chromatography in tandem with mass spectrometry. Recoveries for target analytes were between 35 and 120%, depending on the water matrix. Method detection limits were in the range of 0.5–17 ng L<sup>-1</sup>. The validated method was applied to the determination of perfluorinated compounds in water samples around Ireland. Eight compounds out of fifteen were detected at least in one sample. Measured concentrations were higher in river water than seawater, and drinking water had the lowest levels, although still detectable for a considerable amount of compounds. The most prevalent compounds were PFPeA, PFOA and PFHxA, present in all types of water, and they had the highest concentrations.

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## 1. Introduction

Poly- and perfluoroalkyl substances (PFAS) are a class of thousands of anthropogenic substances that contain an aliphatic fluorinated carbon chain, and that have been in use since the 1950s.<sup>1</sup> The C–F bond in the perfluoroalkyl moiety (–C<sub>n</sub>F<sub>2n+1</sub>, *n* > 1) is highly dipolar and relatively non-polarizable, which imparts the fluorinated compounds with polar hydrophobicity, repelling both water and oil.<sup>2</sup> This has led to widespread industrial and commercial uses, such as construction, chemistry, food production, coatings and paints, surface repellents and textile production.<sup>3–5</sup> As an example, more than 3000 tonnes of PFAS were used just in the Nordic countries.<sup>6</sup> According to the OECD, more than 4700 PFAS compounds have been identified.<sup>7</sup>

Decades of extended global use of these highly persistent (they are often called “forever chemicals”) and mobile compounds have raised the concern about the potential impacts of legacy and new generation fluorinated alternatives on human health and the environment.<sup>8</sup> PFAS can be

bioaccumulated and biomagnified through the food chain.<sup>9–12</sup> PFAS are ubiquitous in the aquatic environment and organisms globally, and have been detected in air, soil, plants and biota.<sup>13–15</sup> Several studies on their toxicity have demonstrated that exposure to PFAS can result in neurotoxicity, hepatotoxicity, developmental toxicity, immunotoxicity, endocrine disruption and carcinogenicity.<sup>16,17</sup>

These studies had led to a great regulatory effort and scrutiny from national and international authorities. PFOS and derivatives have been restricted in Europe for more than 10 years, while PFOA has been banned under the POPs Regulation since 2020.<sup>18</sup> Both PFOA and PFHxS are already included or are being considered to be included in the Stockholm Convention on Persistent Organic Pollutants. In Europe, the recast of the Drinking Water Directive, which took effect on 12 January 2021, includes a limit of 0.5 µg L<sup>-1</sup> for all PFAS, and a limit value of 0.1 µg L<sup>-1</sup> for each individual PFAS.<sup>19</sup> Several EU Member States have set drinking water limits for specific PFAS and for groups of PFAS.<sup>20</sup> PFOS and their derivatives are included as a priority hazardous substance under the EU Water Framework Directive,<sup>21</sup> with a much lower Environmental Quality Standard (AA-EQS) limit value of 0.65 ng L<sup>-1</sup> in inland surface waters and 0.13 ng L<sup>-1</sup> in seawater, while a number of other PFAS are on the REACH list of Substances of Very High Concern (SVHCs).<sup>22</sup> Moreover, several countries have proposed to ban the use of PFAS, except for essential uses under REACH, and it is

<sup>a</sup>DCU Water Institute, School of Chemical Sciences, Dublin City University, Glasnevin, Dublin 9, Ireland. E-mail: belinda.huerta@dcu.ie

<sup>b</sup>Marine Institute, Rinville, Oranmore, Co. Galway, H91 R673, Ireland

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anticipated that the proposal will be submitted to the European Chemical Agency by 15 July 2022.

As a consequence of these regulatory efforts, global manufacturers have started to produce new chemicals in order to substitute long-chain PFAS, such as shorter-chain compounds or with non-fluorinated substances. However, some of these new chemicals have resulted to be equally environmentally persistent and even more mobile in the environment and more difficult to remove from drinking water than their long-chain counterparts, and some have resulted to be toxic as well (*e.g.* GenX).<sup>23,24</sup> In particular, short-chain PFAS accumulate in the environment and have been found to contaminate surface, ground- and drinking water.<sup>25,26</sup>

Therefore, there is an urgent need for continuous monitoring of PFAS in aquatic systems, with special emphasis in drinking water, due to their possible negative impact on human health. In this context, the main objectives of this work were (i) to develop and validate an analytical methodology applicable to drinking water, surface water and seawater; and (ii) to assess the occurrence of 15 PFAS in drinking water in the Dublin area, river water from 4 Irish rivers and seawater from Dublin Bay, with a total of 12 samples analysed. The analytical methodology presented here is an advance for the screening of PFAS in the water cycle in Ireland, and the study of the occurrence of these chemicals will be essential for the implementation of future concentration limits by the regulatory authorities for environmental and human protection.

## 2. Materials & methods

### 2.1. Chemicals and solutions

PFAS substances included in this study were supplied by Wellington Laboratories Inc. (Canda) including: the acids perfluoropentanoic (PFPeA), perfluorohexanoic (PFHxA), perfluorooctanoic (PFOA), perfluorononanoic (PFNA), perfluorodecanoic (PFDA), perfluoroundecanoic (PFUDA), perfluorododecanoic (PFDoA), perfluoropropoxypropanoic (GenX), and perfluorobutanesulfonate (PFBS), perfluorooctanesulfonate (PFOS), perfluorodecanesulfonate (PFDS), perfluoropentylsulfonate (PFPeS), perfluorohexanesulfonate (PFHxS), perfluorooctanesulfonamide (FOSA), perfluorononylsulfonate (PFNS). Internal standards (added before analysis in order to assess and compensate possible losses during sample manipulation and extraction) were purchased from Wellington Laboratories Inc. (Canada) including: [<sup>13</sup>C<sub>8</sub>]-perfluorooctanesulfonate (M8PFOS), [<sup>13</sup>C<sub>2</sub>]-perfluorooctanoic acid (M2PFOA), [<sup>13</sup>C<sub>5</sub>]-perfluorononanoic acid (MPFNNA), and [<sup>13</sup>C<sub>2</sub>]-perfluorohexanoic acid (MPFHxA).

The list of the target analytes, including class, formula and acronyms are listed in Table 1. Individual stock standards and labelled internal standards were prepared in methanol at a concentration of approximately 1000 mg L<sup>-1</sup>. Stock solutions and 50 mg L<sup>-1</sup> mixtures in methanol were stored at -20 °C and diluted to 1 mg L<sup>-1</sup> before each analytical run. Ultrapure water was prepared by a Purelab® Ultra water purification system. Methanol (MeOH) LC-MS grade was obtained from Honeywell (France). Ammonium acetate salt (NH<sub>4</sub>Ac), ammonia (NH<sub>4</sub>OH),

sodium thiosulfate (Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>), acetic acid and acetonitrile (ACN) were obtained from Sigma-Aldrich (Dublin, Ireland).

### 2.2. Sample collection and extraction

The study was conducted in different types of water: (i) drinking water: samples were collected in 4 households in Dublin city; (ii) seawater: samples were collected in from 3 sites in the Dublin Bay area; (iii) surface water: samples were collected from 4 river sections around the western part of Ireland (Rivers Annalee, Liffey, Suir, Nore). Fig. 1 represents the locations where samples were collected during this study.

Water samples were collected in amber polypropylene bottles, previously washed with methanol and rinsed with distilled water, and taken to the laboratory, where pH was adjusted to 3 with acetic acid, and they were kept at 4 °C until extraction if extracted within 24 h; otherwise, they were frozen until analysis. To reduce residual chlorine, 200 µL of 250 mg mL<sup>-1</sup> solution in water of sodium thiosulfate was added to seawater samples.<sup>27</sup> Two types of SPE cartridges (Oasis WAX Cartridges and Oasis HLB, both 500 mg) were conditioned with 5 mL of 0.5% ammonia/methanol (only WAX cartridges), followed by 5 mL of methanol and 5 mL of ultra-pure water (pH 3). Samples (250 mL for river water, 1 L for drinking and marine water) were loaded into the cartridge at a flow rate of 2 mL min<sup>-1</sup>. Cartridges were rinsed with 5 mL of ultra-pure water, and were dried with air for 30 min. Analytes were eluted with 5 mL of methanol followed by 5 mL of 0.5% ammonia/methanol. Eluates were placed under a N<sub>2</sub> current until dryness, and then reconstituted in 1 mL methanol/water (20 : 80). Finally, 5 µL of a 1 mg L<sup>-1</sup> standard mixture containing labelled compounds were added to the extracts as internal standards.

### 2.3. Instrumental analysis

All extracts were analysed by HPLC (Agilent 1260 series Infinity binary pump) coupled to a triple quadrupole mass spectrometer (Agilent 6470), equipped with an electrospray ionization source in negative mode. Ten microliters were injected in an Agilent Zorbax Eclipse Plus C18 column (3.0 × 50 mm; 1.8 µm), set with a guard column of the same materials. Mobile phases were (A): 2 mM formic acid/ammonium formate (pH 3.2) and (B): methanol at a flow rate of 0.4 mL min<sup>-1</sup>. The gradient elution was: 0–1.5 min, 90–45% A; 1.5–7.0 min, 45–10% A; 7.0–8.5 min, 10–0% A; 8.5–10 min, 0% A; 10–11 min return to initial conditions; 11–12 min, equilibration of the column. Column temperature was set at 35 °C. Representative standard chromatogram is shown in Fig. 2.

Compound-dependent mass spectrometric parameters (fragmentor, collision energy) as well as compound-selected reaction monitoring transitions were optimized by direct infusion of individual standard solution of each analyte at 10 ng mL<sup>-1</sup> using the Agilent Compound Optimizer software. A summary of these parameters is presented in Table 2. Source-dependent parameters were determined by flow injection analysis: gas temperature: 230 °C; gas flow: 4 L min<sup>-1</sup>; nebulizer: 15 psi; source temperature, 375 °C; ion spray voltage, 2500; sheath gas flow: 12 L min<sup>-1</sup>. In the cases where two transitions were monitored, the most intense one was used for quantification purposes, whereas the second one was used to



Table 1 Description of target analytes included in the present study

Compound	Acronym	CAS number	Class	Fluorinated C chain	Formula
Perfluorododecanoate	PFDoA	307-55-1	Acid	C12	C <sub>12</sub> HF <sub>23</sub> O <sub>2</sub>
Perfluoroundecanoate	PFUdA	2058-94-8	Acid	C11	C <sub>11</sub> HF <sub>21</sub> O <sub>2</sub>
Perfluorodecanoate	PFDA	335-76-2	Acid	C10	C <sub>10</sub> HF <sub>19</sub> O <sub>2</sub>
Perfluorononanoate	PFNA	375-95-1	Acid	C9	C <sub>9</sub> HF <sub>17</sub> O <sub>2</sub>
Perfluorooctanoate	PFOA	335-67-1	Acid	C8	C <sub>8</sub> HF <sub>15</sub> O <sub>2</sub>
Perfluorohexanoate	PFHxA	307-24-4	Acid	C6	C <sub>6</sub> HF <sub>11</sub> O <sub>2</sub>
Perfluoropropoxypropanoic acid	Gen X	13252-13-6	Acid	C6	C <sub>6</sub> HF <sub>11</sub> O <sub>3</sub>
Perfluoropentanoate	PFPeA	2706-90-3	Acid	C5	C <sub>5</sub> HF <sub>9</sub> O <sub>2</sub>
Perfluorooctanesulfonamide	FOSA	754-91-6	FOSA	C8	C <sub>8</sub> H <sub>2</sub> F <sub>17</sub> NO <sub>2</sub> S
Perfluorodecylsulfonate	PFDS	2806-15-7	Sulfonate	C10	C <sub>10</sub> HF <sub>21</sub> O <sub>3</sub> S
Perfluorononylsulfonate	PFNS	98789-57-2	Sulfonate	C9	C <sub>9</sub> HF <sub>19</sub> O <sub>3</sub> S
Perfluorooctylsulfonate	PFOS	4021-47-0	Sulfonate	C8	C <sub>8</sub> HF <sub>17</sub> O <sub>3</sub> S
Perfluorohexylsulfonate	PFHxS	82382-12-5	Sulfonate	C6	C <sub>6</sub> HF <sub>13</sub> O <sub>3</sub> S
Perfluoropentylsulfonate	PFPeS	630402-22-1	Sulfonate	C5	C <sub>7</sub> HF <sub>13</sub> O <sub>2</sub>
Perfluorobutylsulfonate	PFBS	29420-49-3	Sulfonate	C4	C <sub>4</sub> HF <sub>9</sub> O <sub>3</sub> S
Perfluoro-1-( <sup>13</sup> C <sub>8</sub> ) octanesulfonate	M8PFOS	—	Sulfonate	C8	<sup>13</sup> C <sub>8</sub> F <sub>17</sub> SO <sub>3</sub>
Perfluoro- <i>n</i> -[1,2,3,4,5- <sup>13</sup> C <sub>5</sub> ] nonanoic acid	MPFNA	—	Acid	C9	<sup>13</sup> C <sub>5</sub> <sup>12</sup> C <sub>4</sub> HF <sub>17</sub> O <sub>2</sub>
Perfluoro- <i>n</i> -[1,2- <sup>13</sup> C <sub>2</sub> ] octanoic acid	MPFOA	—	Acid	C8	<sup>13</sup> C <sub>2</sub> <sup>12</sup> C <sub>6</sub> HF <sub>15</sub> O <sub>2</sub>
Perfluoro- <i>n</i> -[1,2- <sup>13</sup> C <sub>2</sub> ] hexanoic acid	MPFHxA	—	Acid	C6	<sup>13</sup> C <sub>2</sub> <sup>12</sup> C <sub>4</sub> HF <sub>11</sub> O <sub>2</sub>

confirm the identity of the target compounds. Instrument control data acquisition and data analysis were carried out using Agilent software.

#### 2.4. Validation

Both MDL and MQL were determined in spiked water samples ( $n = 3$ ) of all matrices considered as the minimum detectable amount of analyte with a signal-to-noise ratio of 3 and 10, respectively. Precision was calculated from five repeated injections of a spiked sample at 1, 50, and 500 ng mL<sup>-1</sup>, and was expressed as the relative standard deviation of the measured concentration.

Calibration curves were generated using linear regression analysis ( $R^2 > 0.99$ ) in the concentration range from 0.01 to 1000 ng mL<sup>-1</sup>. To determine the extraction efficiencies, water samples (drinking, surface and marine water) were spiked with a mixture of the target analytes at 100 ng mL<sup>-1</sup>. Concentrations obtained after the extraction procedure, calculated by internal standard calibration, were compared with the initial spiking levels.

#### 2.5. Background contamination

Careful measures were taken to prevent background levels of PFAS arising from sample manipulation and instrumental

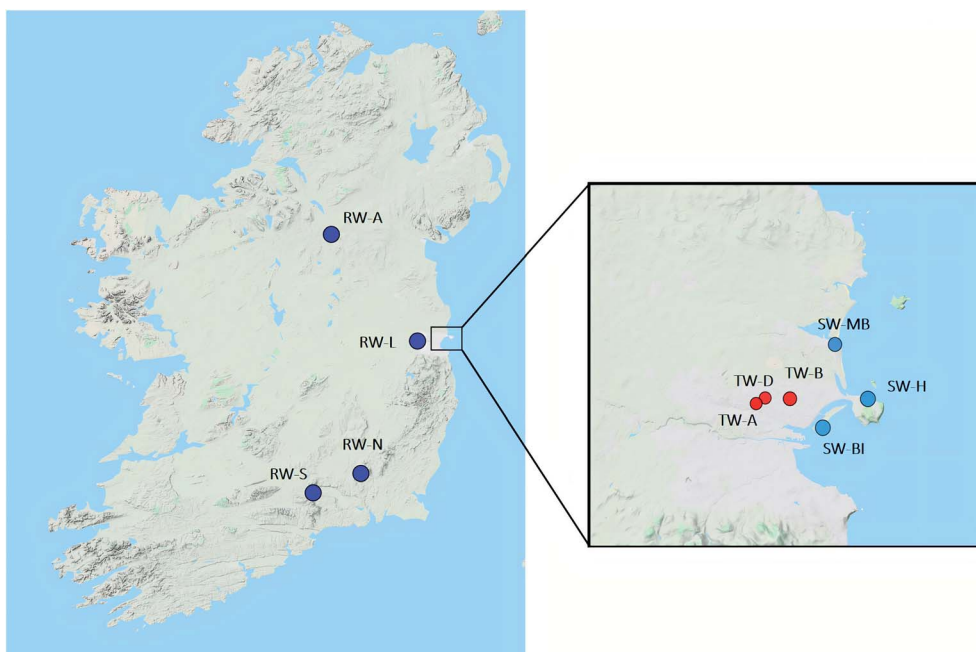


Fig. 1 Samples sites in Ireland. RW: river water. TW: drinking water. SW: seawater.



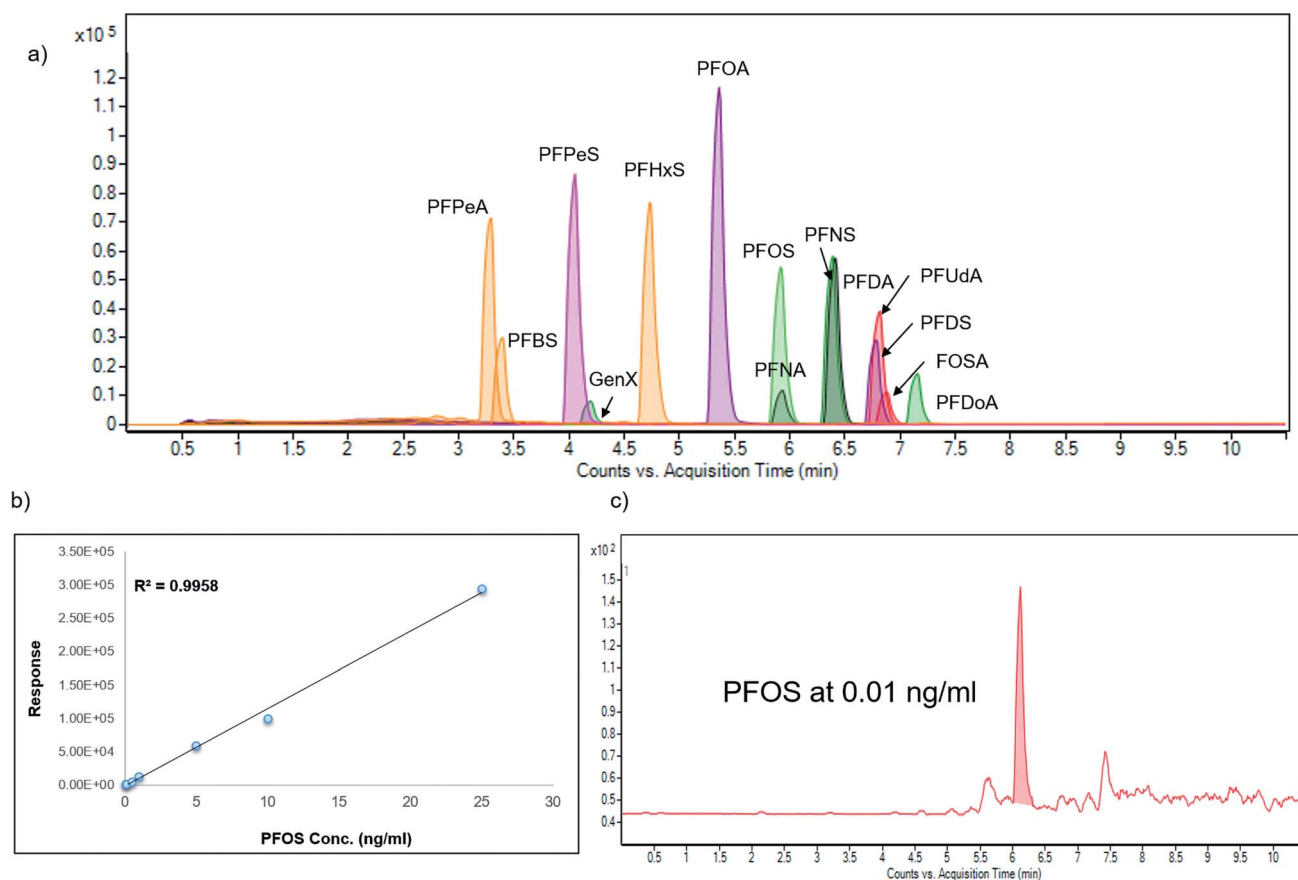


Fig. 2 (a) Compound peaks obtained by injecting a standard mixture at  $100 \text{ ng mL}^{-1}$ ; (b) PFOS calibration curve (levels 0.01 to  $25 \text{ ng mL}^{-1}$ ), with an  $R^2 > 0.99$ ; (c) PFOS chromatogram extracted at a concentration of  $0.01 \text{ ng mL}^{-1}$ .

Table 2 MRM transitions and compound dependant MS parameters for target analytes. For the product ion, second transition was placed between parenthesis. RT: retention time

Compound	Precursor ion	Product ion	RT (min)	Fragmentor	Collision energy (V)
PFDoA	613	519 (219)	7.2	112	8 (20)
PFUdA	563	519 (219)	6.9	112	8 (20)
PFDA	513	469 (269)	6.6	112	8 (12)
PFNA	463	219 (419)	6.1	78	16 (8)
PFOA	413	369 (169)	5.5	78	8 (12)
PFHxA	313	269 (91)	4.1	78	4 (8)
Gen X	285	185 (169)	4.2	78	16 (4)
PFPeA	263	219	3.3	78	4
FOSA	498	78	7.0	180	36
PFDS	599	80 (99)	6.9	180	60 (56)
PFNS	549	80 (99)	6.5	180	56
PFOS	499	80 (99)	6.1	180	44
PFHxS	399	80 (99)	4.9	180	44 (40)
PFPeS	349	80 (99)	4.1	180	40 (36)
PFBS	299	80 (99)	3.4	146	40 (36)
M8PFOS	507	80	6.1	180	52
MPFNA	468	219	6.1	78	16
MPFOA	415	370	5.5	78	8
MPFHxA	315	270	4.1	78	4



analysis. All materials that could potentially contain the target analytes were avoided (Teflon bottles, materials used during SPE, HPLC vials, *etc.*). Polypropylene was used when possible instead of glass to avoid compound retention on the containers. In order to eliminate the sources of contamination from the analytical system all the polytetrafluoroethylene (PTFE) tubing was changed for polyether ether ketone (PEEK) connections.

Procedural blanks were used during sample collection, extraction and treatment prior to instrumental analysis. Ultra-pure water was used with this purpose for optimization and validation, and these samples were analysed to assess the initial concentrations of PFAS. For sample collection, potential PFAS concentration on empty polypropylene bottles was examined by adding 10 mL of methanol and shaking for 10 min. The extract was pre-concentrated to 50  $\mu\text{L}$  and analysed. SPE cartridges were tested for the presence of PFAS through the entire analytical procedure. Cartridges were preconditioned and eluted with 0.5% ammonia/methanol as described in Section 2.2, except no samples were loaded. Solvent blanks (methanol, methanol/water) were also analysed continuously during each run to check for background PFAS levels. Instrumental blanks consisting of initial conditions of the mobile phase were analysed every 5 sample injections to check for carryover and residual PFAS concentration.

## 3. Results & discussion

### 3.1. Method validation

Oasis WAX cartridges were selected for sample extraction, as recoveries were found to be generally higher, and the results were more replicable than with Oasis HLB (see Fig. 3).

Extraction recoveries, method detection and quantification limits were calculated for each target analyte and water matrix (Table 3). Extraction efficiency for all compounds was above

50%, except for PFDS, FOSA, PFuDA and PFoDA (35–48%). Method detection limits (MDLs) were in the range of 0.046–12.62  $\text{ng L}^{-1}$  for drinking water, 0.58–15.15  $\text{ng L}^{-1}$  for river water, and 0.65–17.7  $\text{ng L}^{-1}$  in seawater.

Calibration curves for all compounds prepared in solvent were generated using linear regression analysis ( $R^2 > 0.99$ ) in the range from 0.01  $\text{ng mL}^{-1}$  and 1000  $\text{ng mL}^{-1}$  (Fig. S1†). Instrument accuracy was calculated as the deviation of the measured mean concentration from the real concentration, expressed in percentage from 5 repeated injections of a spiked sample at three different levels (1, 50, 500  $\text{ng mL}^{-1}$ ). Repeated injections were done during the same day (intra-day accuracy) or during 5 consecutive days (inter-day accuracy). At the lowest spiking concentration, values were less than 12.5%, intra-day and 17.5%, inter-day (Table 4).

### 3.2. Comparison of the proposed method with previously published methods

The analytical performance of the present method was compared with other published analytical methods for the determination of PFAS in water samples. Table 5 presents the characteristic data of the present method and the other techniques reported in literature. The method reported here presents comparable MDLs and recoveries to similar methods, in some cases reporting better MDLs and recoveries.

### 3.3. Background contamination

Procedural blanks (ultra-pure water extracted following the same procedure as the samples) were analysed to assess the initial concentrations of PFAS, and in all cases were below the MQLs. Similarly, the residual concentrations analysed from empty polypropylene bottles was considered negligible (results not shown). SPE cartridge blanks (no sample loaded) were

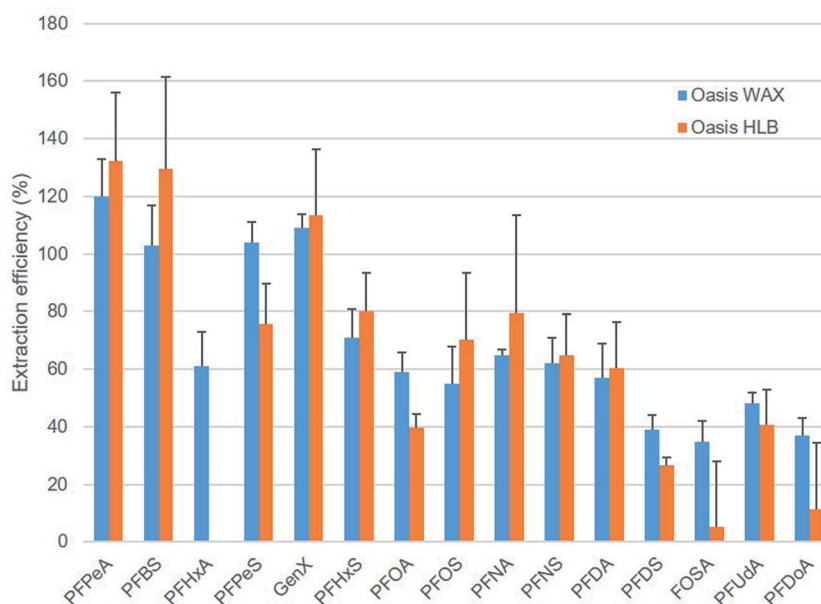


Fig. 3 Extraction efficiencies (%) obtained with two types of SPE cartridges (WAX and HLB) when samples were spiked at 100  $\text{ng L}^{-1}$  ( $n = 3$ ).





**Table 3** Mean percent recoveries ( $n = 3$ ) of 15 PFASs in the three water matrices, and method detection and quantification limits (MDL, MQL) in each water matrix ( $\text{ng L}^{-1}$ )

	Recoveries ( $\pm$ RSD)			MLD ( $\text{ng L}^{-1}$ )			MLQ ( $\text{ng L}^{-1}$ )		
	TW	SW	RW	TW	SW	RW	TW	SW	RW
PFPeA	120 (13)	98 (12)	101 (10)	1.40	1.96	1.68	4.67	6.54	5.61
PFBS	103 (4)	96 (9)	99 (12)	0.48	0.67	0.58	1.60	2.24	1.92
PFHxA	61 (12)	57 (9)	59 (3)	1.78	2.49	2.13	5.92	8.29	7.11
PFPeS	104 (7)	105 (12)	102 (4)	0.75	1.05	0.90	2.49	3.48	2.99
GenX	109 (5)	103 (2)	105 (2)	1.45	2.03	1.74	4.82	6.75	5.79
PFHxS	71 (10)	60 (13)	67 (3)	0.46	0.65	0.55	1.54	2.16	1.85
PFOA	59 (7)	51 (5)	55 (7)	1.27	1.78	1.53	4.24	5.93	5.09
PFOS	55 (13)	59 (9)	55 (12)	0.91	1.28	1.10	3.04	4.26	3.65
PFNA	65 (2)	67 (12)	63 (14)	0.81	1.14	0.98	2.71	3.80	3.26
PFNS	62 (9)	58 (3)	61 (12)	0.66	0.93	0.80	2.21	3.09	2.65
PFDA	57 (12)	55 (9)	59 (10)	0.48	0.67	0.58	1.61	2.25	1.93
PFDS	39 (5)	35 (3)	41 (3)	2.43	3.40	2.91	8.09	11.32	9.70
FOSA	35 (7)	39 (4)	41 (6)	12.62	17.67	15.15	42.07	58.90	50.48
PFUDa	48 (4)	45 (7)	51 (6)	0.91	1.27	1.09	3.02	4.23	3.62
PFDoA	37 (6)	35 (8)	39 (7)	4.11	5.75	4.93	13.69	19.17	16.43

tested for the presence of PFAS through the entire analytical procedure, with no levels detected above the MQLs for any compound. Instrumental blanks (methanol/water) were analysed every 5 sample injections, and no peaks were detected above the MDLs.

### 3.4. Environmental samples

A total of 12 water samples were analysed for method validation purposes, including a procedural blank (TW-C) and a solvent blank (TW-M). Results obtained are shown in Table 6 (data adjusted with recoveries). Fig. 4 shows the relative quantities of target analytes in each water sample. Out of the 15 compounds in the developed method, 8 were detected at least once in river water (Fig. S2†). These samples contained the highest PFAS

concentrations, with a maximum detected of  $424 \text{ ng L}^{-1}$  for PFPeA, followed by PFOA (approx  $200 \text{ ng L}^{-1}$ ), similarly to other studies carried out in surface waters, where PFAS levels have been found in the low  $\text{ng L}^{-1}$  range in rivers of urban catchments.<sup>31–33</sup> In seawater samples, 7 compounds were detected, but only 3 were above MQLs. The highest concentrations were around 45 and  $13 \text{ ng L}^{-1}$  for the compounds PFBS and PFPeA, respectively, an order of magnitude lower than in river water, comparable to similar studies in ocean water. However, some studies reported concentrations at several orders of magnitude greater in highly polluted areas.<sup>27,34,35</sup> In drinking water samples, only 5 compounds were detected with values  $>$  MQLs, in the range of  $4\text{--}7 \text{ ng L}^{-1}$ , with very few exceptions. The most prevalent compounds were PFPeA and PFHxA, present in all

**Table 4** Intra-day and inter-day accuracy (expressed as % RSD) calculated at three spiking levels for 15 PFAS

	Accuracy (% RSD)					
	Intra-day			Inter-day		
	1 $\text{ng mL}^{-1}$	50 $\text{ng mL}^{-1}$	500 $\text{ng mL}^{-1}$	1 $\text{ng mL}^{-1}$	50 $\text{ng mL}^{-1}$	500 $\text{ng mL}^{-1}$
PFPeA	1.4	0.9	2.4	17.5	2.4	10.9
PFBS	6.9	0.9	1.9	0.5	4.6	0.8
PFHxA	2.9	15.5	6.7	1.7	2.6	5.5
PFPeS	2.0	1.5	1.5	0.8	1.3	7.3
GenX	12.4	1.8	1.5	7.2	5.1	0.0
PFHxS	0.7	1.2	4.3	1.2	0.8	7.7
PFOA	0.7	1.2	4.3	4.2	2.2	9.4
PFOS	4.0	1.0	2.2	1.7	1.7	2.7
PFNA	2.1	0.4	1.1	2.2	0.9	11.1
PFNS	5.1	2.4	10.2	3.0	1.7	11.0
PFDA	1.2	0.4	10.3	2.5	1.0	2.0
PFDS	2.9	2.5	1.3	3.7	3.5	3.8
FOSA	2.6	0.4	1.1	9.2	10.0	7.6
PFUDa	11.7	1.3	2.5	5.4	5.0	10.4
PFDoA	5.3	1.3	2.2	3.1	2.2	7.5



Table 5 Comparison of the present method with other analytical methods for the determination of PFAS in water samples

Reference	Compounds	Extraction	Detection	Sample	Linearity ( $R^2$ )	MDLs (ng L <sup>-1</sup> )	Recovery (%)	Accuracy <sup>a</sup>
Present method	15	SPE	HPLC MS/MS	Tap, river, seawater	>0.99	0.5–17.7	35–120	1–12%
(Liang <i>et al.</i> , 2020) <sup>28</sup>	21	SUPRAS-based microextraction	HPLC-Orbitrap	Tap, river	>0.99	10–80	72–118	1–11%
(Pico <i>et al.</i> , 2012) <sup>29</sup>	10	SPE	HPLC-MS/MS	Surface water	>0.99	0.02–1.5	55–85	Not reported
(Gebbinck <i>et al.</i> , 2017) <sup>26</sup>	14	SPE	UPLC-MS/MS	Tap, river	>0.99	0.01–4	81–115	Not reported
(Ali <i>et al.</i> , 2021) <sup>30</sup>	13	SPE	HPLC MS/MS	Seawater	>0.99	0.04–4.1	17–117	Up to 10%

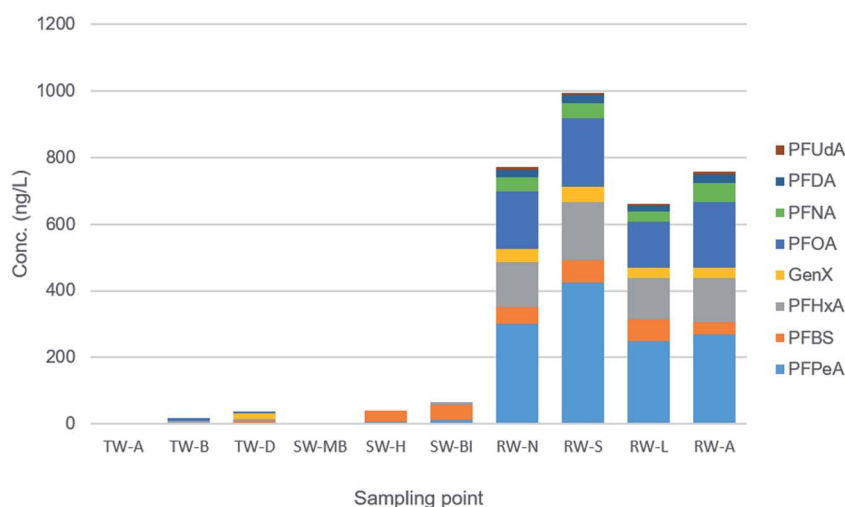
<sup>a</sup> Intra-day precision at the lowest spiking level.

Table 6 Concentration (ng L<sup>-1</sup>) of detected PFAS in environmental samples. BLK 1 represents the average results of the procedural blanks (ultra-pure water,  $n = 3$ ). BLK 2 represents the SPE cartridge blank average results ( $n = 3$ ). TW: drinking water. SW: seawater. RW: river water

Matrix	PFPeA	PFBS	PFHxA	GenX	PFOA	PFNA	PFDA	PFUdA
BLK 1	Blank H <sub>2</sub> O	<MLQ	<MLQ	<MLQ	<MQL	<MLQ	<MLQ	<MDL
BLK 2	Blank SPE	<MLD	<MDL	<MDL	<MQL	<MDL	<MQL	<MDL
TW-A	Drinking water	<MLQ	<MDL	<MDL	<MQL	5.3	<MQL	<MDL
TW-B	Drinking water	<MLQ	<MDL	10.7	<MQL	7.3	<MQL	<MDL
TW-D	Drinking water	5.3	4.1	6.6	15.6	6.8	<MQL	<MDL
SW-MB	Seawater	<MLQ	<MQL	<MQL	<MQL	<MQL	<MQL	<MDL
SW-H	Seawater	6.7	34.0	<MQL	<MQL	<MQL	<MDL	<MDL
SW-BI	Seawater	13.5	45.3	7.0	<MQL	<MQL	<MQL	<MDL
RW-N	River water	301.2	49.5	134.6	39.1	172.9	43.0	22.2
RW-S	River water	424.1	68.9	173.3	44.7	205.4	46.5	24.1
RW-L	River water	250.2	64.0	122.7	33.4	138.2	30.8	15.8
RW-A	River water	269.5	35.2	132.6	32.0	197.5	56.4	23.7

types of water, and they had the highest concentrations. Literature shows that drinking water PFAS levels are usually low (pg L<sup>-1</sup> range), but some studies have found them up to the µg L<sup>-1</sup> level.<sup>36,37</sup> As expected, concentrations were much higher in river water than seawater, and drinking water had the lowest levels,

although still detectable for a considerable amount of compounds. This screening provides a valuable snapshot of what the situation is in Irish waters, but a more extensive monitoring campaign is required to understand the impact of

Fig. 4 Concentration (ng L<sup>-1</sup>) of detected pfas in environmental samples. TW: drinking water. SW: seawater. RW: river water.

seasonality and land use on the occurrence of these compounds in water systems.

According to European legislation,<sup>18</sup> levels of PFOS should be lower than 0.65 ng L<sup>-1</sup> in surface water and lower than 0.13 ng L<sup>-1</sup> in seawater. In this study, PFOS was not detected in any of the samples analysed. Quantification of PFAS in seawater at the required protection level and associated performance criteria for marine waters is particularly challenging. The detection limits achieved in this method (*e.g.*, for PFOS in seawater 1.28 ng L<sup>-1</sup>) would allow for the method to be applied for ultra-trace screening of marine waters. For drinking water, the addition of all concentrations reached a maximum of 65 ng L<sup>-1</sup>, well below the 500 ng L<sup>-1</sup> limit by the Drinking Water Directive.<sup>21</sup> More concerning is the occurrence of GenX in surface waters at concentrations around 40 ng L<sup>-1</sup>. Although GenX is not currently regulated, this compound has demonstrated to be equally toxic than some of the legacy PFAS,<sup>24,38</sup> and the US EPA has calculated that chronic toxicity for human consumption could be ten times greater than PFOS.<sup>39</sup>

## 4. Conclusions

While there are several LC-MS/MS based analytical methods for PFAS analysis reported in the literature,<sup>13,26,27,36,40</sup> reliable analytical methods for trace determination of legacy and emerging PFAS in varied media, suitable for routine environmental risk assessment, is highly desirable. This study developed a simple, rapid, and efficient analytical method for the determination of 15 PFASs, including legacy and novel compounds, some of them heavily regulated, in different types of water: seawater, river and drinking water. The developed method allows the identification of short and long chain compounds, which in many cases it is linked to the type of application/source of each compound. The final procedure consisted of pre-concentration of water samples with anionic exchange solid-phase extraction, and the analysis was performed by HPLC-MS/MS, which provided the necessary sensitivity and selectivity for these contaminants. The method was applied to assess the occurrence of these contaminants in Ireland. The assessment of PFAS in real samples showed that PFPeA, PFOA, PFBS, and PFHxA were present in all types of water, and they occurred at the highest concentrations in all matrices. The highest concentrations were found in surface river waters, with a maximum value detected of 424 ng L<sup>-1</sup> for PFPeA in the River Suir (Ireland). For drinking water, the levels detected were below the limits stated by the Drinking Water Directive.<sup>19</sup>

## Conflicts of interest

There are no conflicts to declare.

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