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Pharmaceutical compounds in urban drinking waters of Ecuador

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Introduction: Emerging contaminants (ECs) are substances with widely diverse chemical structures that may pose a risk to the environment and human beings. The limited scope of water treatment facilities, particularly in low and middle-income countries, allows ECs to be continuously introduced to the environment and become part of the cycle again for potabilization. In this work, we study for the first time the presence of these compounds in the drinking water of five Ecuadorian cities.

Methods: The compounds of interest were mainly pharmaceutical substances commonly present in prescription and over-the-counter medicines, along with caffeine, a known coadjuvant in some of these preparations. Samples were collected from Quito, Guayaquil, Cuenca, Ibarra, and Esmeraldas, considering each city's distribution systems, and, after solid-phase extraction, analyzed by LC-MS/MS ESI+.

Results and discussion: Results showed a high occurrence of caffeine, the only analyte present in all cities, with concentrations ranging from <6.35 to 201 ngL⁻¹ and an occurrence from 11% in Quito to 77% in Cuenca. The highest median was found in Cuenca, followed by Esmeraldas. Our observations regarding concentrations are comparable to other studies around the globe. Although in other cities, some pharmaceuticals appeared at levels below our detection limits. These findings highlight the ubiquitous nature of emerging contaminants while pointing out the need for regulatory frameworks that facilitate the implementation of treatment technologies at the source and wastewater level. These actions will safeguard public and environmental health in the long term.

KEYWORDS

emerging contaminants, drinking water, Ecuador, pharmaceuticals, caffeine

1 Introduction

Emerging contaminants (ECs) are naturally occurring or synthetic compounds that are not commonly monitored in the environment and pose potential risks to human and environmental health (Naidu et al., 2016). There are several types of ECs, including pharmaceutical products (e.g., antibiotics, painkillers, anti-inflammatories) and associated substances. Medicaments reach drinking water systems through the introduction of wastewater effluents into rivers, which serve as sources for water treatment facilities. Current technologies employed in these plants are not effective in removing pharmaceutical compounds before distribution (Furlong et al., 2017).

Over the past 2 decades, emerging compounds have garnered significant attention from the scientific community due to their impact on human health and ecosystems (Ramírez-Malule et al., 2020). While some compounds may undergo changes when exposed to the environment, their continuous introduction offsets such effects. Studies have also demonstrated the influence of socioeconomic factors in the occurrence of ECs, such as their higher concentrations in regions with higher values of gross domestic product per capita and human development index (Santos et al., 2020). Furthermore, the constant consumption of pharmaceuticals can lead to bacterial resistance (Tell et al., 2019) and their physicochemical properties, such as high water solubility and low biodegradability (Gil, Soto, Usma and Gutiérrez, 2013), can result in bioaccumulation increasing the risk associated with these substances (Du et al., 2014; Muir et al., 2017). Additionally, there is evidence of environmental risks, particularly to marine fauna. Li and Lin (2015) found that a 2-h exposure to a mix of pharmaceutical compounds including sulfamethoxazole, caffeine, diclofenac, and acetaminophen, led to increased mortality and abnormal behaviors in fish. The concentrations used in the study (3.9 mgL⁻¹ each, previously exposed to sunlight) were similar to those found in local hospital raw wastewater.

Most of the investigations on ECs have been conducted in North America and Europe, where monitoring of pharmaceuticals, perfluorinated chemicals, estrogenic hormones, detergents, microplastics, and inorganic elements in surface and drinking waters has been carried out (Furlong et al., 2017; Sousa et al., 2018). The United States, China, Spain, Italy, and Canada have emerged as the top five nations with the most published works in this field (Ramírez-Malule et al., 2020). However, limited research has focused on the presence of ECs in drinking water in Latin America, with findings reported only in Brazil, Colombia, Venezuela, and Chile, representing less than one-third of the countries in the region (Peña-Guzmán et al., 2019). Besides the limited information, the lack of regulation and monitoring, along with the poor water treatment and purification processes result in an increased presence of these compounds in all bodies of water (Vargas-Berrones et al., 2020). Pinos-Velez et al. (2019) made a comprehensive review of ECs occurrence in Latin America and reported that caffeine was the most prevalent and concentrated compound, although triclosan, cocaine, non-ylphenol, bisphenol A, atrazine, ibuprofen, among others, were also detected in estuaries and drinking water.

In Ecuador, less than 25% of wastewater receives treatment before being discharged into rivers and seas, leading cities like Guayaquil and Esmeraldas to receive industrial and domestic residual effluents in their drinking water sources (Pinos Velez et al., 2019). Studies have sulfamethoxazole, found caffeine. venlafaxine. o-desmethylvenlafaxine, and some steroidal estrogens throughout the San Pedro, Guayllabamba, and Esmeraldas River basins (Voloshenko-Rossin et al., 2015), as well as caffeine, paracetamol, and trimethoprim in the north of the Ecuadorian Amazon (Capparelli et al., 2021). Since ECs have already been detected in water bodies in Ecuador, and considering that most municipalities directly discharge wastewater into various river basins, it becomes imperative to determine whether these compounds are reaching the drinking water systems of urban areas. We hypothesize that they are and that this information may shed light on the environmental impact of ECs on the national ecosystem. Therefore, our research aims to answer the following questions: Are there emerging compounds present in the drinking water of five representative cities in Ecuador, namely: Quito, Guayaquil, Cuenca, Esmeraldas, and Ibarra? At what concentration are they found?

2 Materials and methods

2.1 Study area

As detailed in Table 1, the three most populated cities of Ecuador were selected for this study, namely: Quito, Guayaquil, and Cuenca. We also included two smaller cities that are within the top 10% in terms of population. Esmeraldas, a small coastal city, has previous reports of emerging contaminants in the namesake river (Voloshenko-Rossin et al., 2015) which is also the source for water potabilization and in other water bodies that surround the city (Cipriani-Ávila et al., 2023). Besides, the province of Esmeraldas, where the city is located, has low indexes of potabilization and wastewater treatment. In contrast, Ibarra is the capital city of Imbabura, one of the provinces with most potabilization and wastewater treatment facilities, as well as one with the highest compliance with the technical norm NTE 1 108: 2011, which establishes drinking water requirements (INEC, 2016).

The sampling points in each city, detailed in Figure 1, were chosen based on their respective drinking water production systems, their distribution networks and their main parishes. In Quito, for each of the water distribution subsystems (Bellavista, Puengasí, El Troje, Northwest Treatment Plant) three neighborhoods were randomly selected, within each neighborhood three sampling points were picked out. In the case of Guayaquil, the city was divided into northern, central, and southern sectors. Samples were taken from five in each sector; Alborada neighborhoods and Urdesa neighborhoods were considered twice, as they cover two sectors of the city, Cuenca's drinking water system is divided into three main subsystems named after the rivers from which the water is drawn: Tomebamba, Machángara, and Yanuncay. For each of these, sampling points were selected in their distribution zones in Cuenca city.

In Esmeraldas 2-3 random sampling points were selected per area, namely: south, center, high neighborhoods, low neighborhoods, Las Piedras and Tachina; these areas were established according to the water distribution system and the regularity of supply. In Ibarra, we worked in the five urban parishes (El Sagrario, San Francisco, Alpachaca, Priorato and Caranqui), where three sampling points

TABLE 1 Main characteristics of the Ecuadorian cities that were part of this study.

	Quito	Guayaquil	Cuenca	Esmeraldas	Ibarra
Population	2.239.191ª	2.350.915ª	505.585ª	189.504ª	181.175 ^a
Elevation (m)	3058 ^b	21 ^b	2525 ^b	115 ^b	3140 ^b
Average annual temperature (°C)	12.2 ^b	16.2 ^b	12.4 ^c	25.7 ^b	10.7 ^b
Average annual precipitation (mm)	125.0 ^b	84.9 ^b	126.5°	206.3 ^b	109.5 ^b
Water production company	Empresa Pública Metropolitana de Agua Potable y Saneamiento Quito (2023)	Interagua (2023)	Empresa de Telecomunicaciones, Agua Potable, Alcantarillado y Saneamiento de Cuenca [ETAPA EP] (2023)	Empresa Pública Mancomunada de Agua Potable y Saneamiento de Esmeraldas (2021)	EMAPA-I
Main water sources	Antisana volcano zone; Atacazo and Lloa water subsystems; Pita River ^d	Daule River ^d	Tomebamba, Machángara Yanuncay, Culebrillas, Irquis Rivers ^d	Esmeraldas River ⁴	Yuyucocha, Guaraczapas, Palestina natural springs. Yuyucocha underground wells ^e

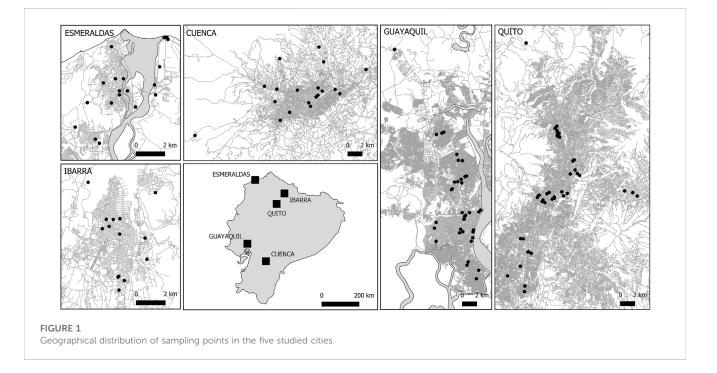
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^aINEC (National Institute of Statistics and Census).

^bInstituto Nacional de Meteorología e Hidrología (2023). ^cClimate-Data.org (2023)

^dOfficial websites of water production companies.

^eJapanese International Cooperation Agency (2005).



were chosen. At each sampling point, one water sample was taken, adding to a total of: 45 samples in Quito, 42 in Guayaquil, 15 in Cuenca, 15 in Ibarra and 16 in Esmeraldas.

2.2 Sampling and sample preparation

Sampling was carried out with the methodology described by the International Standardization Organization NTC-ISO 5667–1:2008.

Five hundred milliliters of each sample were placed in an amber glass bottle with screw cap and subsequently stored at 4°C until processed at the lab. The time between sampling and processing was not longer than 24 h in all cases.

Samples underwent the methodology by Glassmeyer et al. (2017). Analytes were separated from the water matrix by solid-phase extraction (SPE) using a vacuum pump (Millipore, WP6111560), a manifold ($27 \times 17 \times 9.5$ cm), and OASIS HLB cartridges (Waters, 200 mg, 6 ml). To condition the cartridges, they

	Caffeine	Sodium diclofenac	Acetaminophen	Sulfame- thoxazole	Trimethoprim
Retention time (min)	3.23	7.47	2.64	4.59	3.43
Precursor ion (m/z)	195	295.8	152	253.9	291.2
Product ions (m/z)	138, 110	214.7, 249.9, 277.6	109.9, 65.1, 93	92, 155.8	230
Limit of detection (ngL ⁻¹)	6.35	12.5	6.25	3.13	12.5
Limit of quantitation (ngL ⁻¹)	12.5	18.8	12.5	6.25	18.8
Recovery (%)	83	85	80	77	90

TABLE 2 Analytical method performance and relevant parameters for each analyzed compound.

were treated with 6 ml of reagent water after 4 ml of methanol at a flow rate of 10 ml/min. After eluting 500 ml of sample, the cartridges were dried under vacuum for 10 min before analytes were eluted into glass tubes using 6 ml of methanol. Using nitrogen, extracts were concentrated to dryness and reconstituted to 0.5 ml with methanol, filtered through a 0.22 μ m filter, and then placed into vials for chromatographic analysis. As part of measures to assure the quality of the results, a field blank was collected and analyzed to assess that sampling and extraction procedures were performed correctly. In addition, a spiked sample of the mean concentration level was used to gauge the recovery rates of each component (Table 2). All analytical standards had a purity >96.8% and were obtained from Supelco or Sigma-Aldrich.

2.3 Instrumental analysis

Five of the most consumed pharmaceutical compounds in Ecuador were selected for this study, namely: caffeine (stimulant), acetaminophen (analgesic), sodium diclofenac (anti-inflammatory), sulfamethoxazole (antibiotic), and trimethoprim (antibiotic). The information was obtained by request of the authors in May 2019 from the National Undersecretariat of Public Health Governance (Subsecretaría Nacional de Gobernanza de la Salud Pública del Ecuador, 2019).

These compounds were analyzed by UPLC-Qtof (Waters Model I-class liquid chromatograph coupled to a Waters Xevo G2 QTOF) under the conditions described in Cipriani Avila et al. (2023). Briefly, acetonitrile/formic acid in a 99.9%/0.1% ratio (solvent B) and 0.1% formic acid in water (solvent A) were the solvents utilized as the mobile phase; the elution gradient was: 5% B over 1 min, 5%-100% B over 9 min, 100%-5% B over 2 min, and then a column reequilibration at 5% B during 3 min. A C18 column (Waters Acquity BEH 1.7 µm, 100 mm 2.1 mm i. d.) operating at 25°C was used with a flow rate of 0.3 ml min⁻¹. For the electrospray ionization (ESI-MS), a positive mode with a capillary voltage of 0.5 kV, $30 \text{ L} \text{ h}^{-1}$ cone gas flow, $900 L h^{-1}$ desolvation gas flow, $120^{\circ}C$ source temperature, 450°C desolvation temperature with sampling cone and source compensation at 40 and 80 V, were employed. Further fractionation (MS/MS) was conducted with ramp collision energy 20-30 eV. The range of m/z was 50-1,000 Da. Because the acquisition method provides complete information, the analytes were quantified using the precursor or product ion with the highest intensity. The method's performance was determined according to Eurolab España and Morillas (2016).

2.4 Statistical analyses

For the statistical analysis, the software R version 4.2.1 (RRID: SCR_001905) was used with the R-Studio interface. Non-parametric Kruskal–Wallis statistics were performed to determine significant differences in caffeine content in water by city at a significance level of 0.05. Once statistically significant differences were found, the pairwise Wilcox test was applied.

3 Results

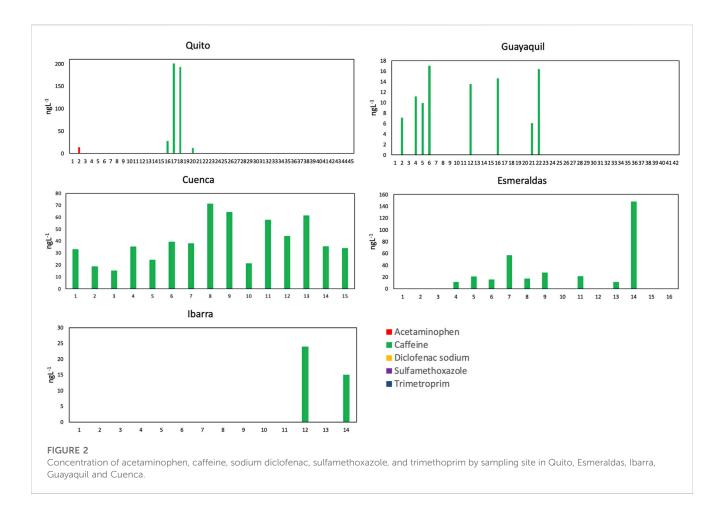
Table 2 indicates the main figures of merit of the analytical method employed for the emerging contaminants in this study. The highest sensitivity was obtained for sulfamethoxazole, although the recovery was lower compared to the other CEs. Trimethoprim showed the highest recovery among the five tested compounds but also the highest limits of detection and quantitation.

Figure 2 shows the results of the five studied compounds in each city. Caffeine was found in all cities with an overall concentration range of $1.4-201 \text{ ngL}^{-1}$. It is also observed that acetaminophen was only found in site 20 (Bellavista) of Quito with 13.2 ngL^{-1} (occurrence 2.2%).

In Table 3 it is observed that the occurrence of caffeine varies from 11% in Guayaquil to 100% in Cuenca. The highest and lowest concentrations were found in Quito with 201 and 1.4 ngL^{-1} , respectively. In the cities with higher occurrence, Cuenca and Esmeraldas, where more than half of the samples contained caffeine, the variation in concentration ranged from 11.6 to 148 ngL⁻¹. Sodium diclofenac, sulfamethoxazole, and trimethoprim were not found in any sample.

In Figure 3 the notch boxplot of caffeine per city is presented. The highest median was found in Cuenca, approximately 36 ngL⁻¹ with 95% of confidence. Esmeraldas, Guayaquil, Ibarra, and Quito show outliers. Kruskal–Wallis found statistically significant differences between medians (chi-squared = 62.3, p < 0.001). Through the pairwise Wilcox test for comparison of median, differences were found between Cuenca and all the other cities with Cuenca with the highest median. Also, differences were found between Esmeraldas with Guayaquil and Quito where Esmeraldas presented the highest median. Moreover, differences were found between Ibarra with Guayaquil and Quito where Ibarra has the highest median.

To contextualize the results of our research, in Table 4 we provide data from similar studies conducted in drinking water



		Caffeine				
	Occ., %	Median*, (ngL ⁻¹)	Min-Max, (ngL ⁻¹)	Mean, (ngL^{-1})	STD, (ngL ⁻¹)	
Quito	11.1	27.5	1.4-201	87.05	100.9	
Guayaquil	19.04	12.35	6.1-17	11.98	4.09	
Cuenca	100	35.75	15.5–71.5	39.75	17.15	
Esmeraldas	56.3	20.9	11.6–147.9	36.88	48.88	
Ibarra	13.33	19.5	15-24	19.5	6.36	

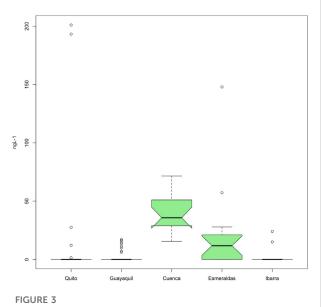
BLD: below the limit of detection.

*Calculated using concentrations greater than the LOD, of the compound.

(after potabilization process) around the globe. In all of these cases, the methodology was similar to ours, although some minor differences in certain references are noted later in the discussion. The number of analyzed samples has a wide range, from 8 to 179; this study is the one that presents the higher value albeit comprising a broader geographical distribution. Regarding the analytes, it can be seen that caffeine and sulfamethoxazole are reported more frequently, followed by diclofenac and acetaminophen; except for one, the concentrations are in the same order of magnitude as ours but in some cases, the values are below the detection limits of our study.

4 Discussion

This is the first study to show the concentration ranges of highoccurrence contaminants in the drinking water supplies of Ecuadorian cities: Quito, Guayaquil, Cuenca, Esmeraldas, and Ibarra. These compounds were determined using SPE followed by UPLC/ESI-MS; the method provided satisfactory analytical performance for the study of all five substances. The findings revealed the presence of two of the five compounds investigated, with caffeine having the highest occurrence, ranging from 11% of samples in Quito to 77% in Cuenca. Concentrations were



Notch boxplot of caffeine $({\rm ngL^{-1}})$ for Quito, Esmeraldas, Ibarra, Guayaquil, and Cuenca.

comparable to those reported in other cities worldwide, hence adding to the evidence that there is an increasing need for regulation and management of these compounds in bodies of water.

According to Figure 2 and Table 3, caffeine was the emerging contaminant with the highest occurrence in drinking water samples, being found in 11% of the Quito samples, 19% in those of Guayaquil, 13% of Ibarra's, 56% of Esmeraldas', and 100% of Cuenca's. However, as shown in Figure 3, despite the low occurrence, Quito had one of the highest variations in concentrations, as caffeine ranged from 1.4 ngL^{-1} to 201 ngL⁻¹. When comparing with other countries of the continent, these values are intermediate between those reported for Canada and Brazil; in them, maximum values of 120 ngL⁻¹ and 2769 ngL⁻¹ were found (Machado et al., 2016; Pulicharla et al., 2021). Given the high frequency of caffeine-containing samples, there were significant

differences between the results for Cuenca and the other four cities. It is not possible to pinpoint the causes for the increased caffeine concentration observed in Cuenca; however, we think that the tourism and cattle-raising activities near the catchment waters at the Cajas National Park might be a key influence. Lodging and restaurants in the Cajas National Park do not have a wastewater treatment plant or garbage disposal systems, which is why wastewater infiltrations and a clandestine garbage dump have been detected (Beltran, 2023). The drinking water in Esmeraldas showed a narrower distribution compared to Cuenca while presenting a higher overall concentration than Quito, Guayaquil, and Ibarra. This may be explained by the quality of the source water. Esmeraldas River receives the wastewater, treated and not, from the cities in the mountain region, being Quito the biggest one. As previously reported in the study by Voloshenko-Rossin et al. (2015), caffeine was among the emerging contaminants detected in this river, whose waters are the intake for the potabilization process in the city.

Caffeine is a commonly used psychoactive ingredient in free trade medicine and daily beverages, and although a large part of the intake is metabolized, it is accepted as an indicator of anthropogenic contributions to water bodies (Daneshvar et al., 2012; Li et al., 2020). The presence of this compound in wastewater in Ecuador has been reported by Capparelli et al. (2021) and Arcentales-Ríos et al. (2022); the latter group of researchers also demonstrate that only 50% of caffeine concentration is decreased with traditional technologies employed in Ecuador. In the best-case scenario, this treated water is released into the environment and eventually is used as a source for drinking water treatment. In other cases, untreated wastewater, rich in emerging contaminants, is directly discharged into these sources. Indeed, treated and untreated wastewater are the largest source of emerging pollutants in the environment since these are discharged to surface waters, distributing them in natural environments where they endanger flora and fauna (Pinos-Vélez et al., 2019). Currently, conventional primary and secondary treatments such as activated sludge, up-flow anaerobic sludge blanket reactor, stabilization ponds, wetlands, and trickling filters only contribute to a small portion of caffeine removal (Zhou et al., 2010; Martín et al., 2012). Organic matter removal mechanisms are unsuitable for compounds with more stable chemical structures like caffeine and other

	Caffeine	Sodium diclofenac	Acetaminophen	Sulfame- thoxazole	Trimethoprim
Ecuador $n = 179$ (this study)	201	<12.5	13.2	<13.3	<12.5
Brazil $n = 14$ Sodré and Sampaio, (2020)	7.8	6.03		5.0	
Brazil n= 12 de Oliveira et al., (2019)			1.6		
Brazil $n = 100$ Machado et al., (2016)	2769				
Canada n = 25 Pulicharla et al., (2021)	120		210	6.4	
China n = 36 Ben et al., (2020)				0.98	5.2
Malaysia $n = 155$ Wee et al., (2020)	5.33	21.39		0.90	
Taiwan $n = 18$ Pai et al., (2020)	263.2				
Hungary n = 108 Kondor et al., (2021)	38.41	4.2			

TABLE 4 Comparison of pharmaceutical compounds concentrations (ngL⁻¹) in drinking waters around the world. Only the highest reported value is shown.

emerging pollutants (Arcentales-Ríos et al., 2022). For this reason, the incorporation of provisional or definitive ternary processes (like adsorption or advanced oxidation processes) into wastewater treatment plants is crucial to eliminate emerging contaminants (Morin-Crini et al., 2022).

Studies show that pharmaceuticals are also inefficiently removed in conventional potabilization treatments (Padhye et al., 2014; McKie et al., 2016), This is evidenced in the results of various studies, such as those presented in Table 4. Our results back up these findings, as caffeine was found in the drinking water of all the studied cities. Although the caffeine concentrations we found in drinking waters do not represent a risk to human health, they put in evidence the ineffectiveness of water treatment plants to eliminate emerging contaminants, both in wastewater and drinking water processing. Considering that with the current lifestyle, introduction rates, as well as EC diversity, tend to increase, it is important to take action to stop the introduction of these compounds into our water sources. For instance, alternative procedures have been proposed that allow the complete elimination of emerging compounds in drinking water, such as activated carbon processes, advanced oxidation processes, ozonation, among others (Kim et al., 2007; Yang et al., 2017; OECD, 2019).

The occurrence of acetaminophen was very low in Ecuadorian cities. Only one sample in Quito showed a quantifiable value for this compound with a concentration of 13.20 ngL⁻¹. As detailed in Table 4, higher values have been reported by Pulicharla et al. (2021). The results found in our study may be due to the transformation of paracetamol during the chlorination of water in the treatment processes, where the main reaction products are hydroquinone and two types of chlorinated compounds: monochlorinated acetaminophen and dichlorinated acetaminophen (Cao et al., 2016). It is also possible that the concentrations in the analyzed samples were similar to those reported by De Oliveira et al. (2019), thus below the detection limit of our analytical method. As for the health impact of acetaminophen, the Minnesota Department of Health recommends a maximum value in drinking water of 200 ppb, equivalent to $200 \ \mu g L^{-1}$, pointing out that the liver might be the organ most prone to damage by chronic exposure (Minnesota Health Department, 2014). Our results are more than an order of magnitude below this threshold.

Although sodium diclofenac, sulfamethoxazole, and trimethoprim were not determined in the drinking water samples from the Ecuadorian cities, some previous studies show their presence in this matrix in other territories. Kondor et al. (2021) indicated a maximum concentration of 4.2 ng sodium diclofenac per liter of drinking water in Hungary. Sulfamethoxazole was frequently found in the samples of a study carried out in the United States where 51 emerging compounds were analyzed (Benotti et al., 2009), as well as in the drinking water of cities in Brazil and Spain with values less than 10 ngL⁻¹ (Gros et al., 2012; Monteiro et al., 2017). As with acetaminophen, this compound may have been chemically transformed during potabilization, due to the high reactivity of sulfonamides with chlorine or chlorine dioxide (Huber et al., 2005; Gaffney et al., 2016). The absence of trimethoprim in the samples can be explained by a lack of sensitivity of the selected analytical methodology, as detailed below.

The publications by Machado et al. (2016) and Gros et al. (2012) show results of LC-MS/MS methods for several emerging compounds, obtaining method LODs of 1, 0.3, 0.8, 0.1, and 0.1 ngL⁻¹ for caffeine, sodium diclofenac, acetaminophen, sulfamethoxazole, and trimethoprim, respectively. Wee et al. (2020) also report lower method detection limits. In contrast, as detailed in Table 2, the LOD values obtained for this research were higher for all analytes. These differences can be attributed to method development. For instance, we focused on sensitivity rather than resolution for spectral scanning optimization. On the other hand, our analyses were conducted in positive ionization mode, whereas, Wee et al. (2020) and Gros et al. (2012) use ionization in both positive and negative modes, probably because certain compounds exhibit higher ionization in the latter, thereby yielding greater analysis sensitivity and, consequently, lower detection limits. Furthermore, Gros et al. (2012) applied ESI- along with acetonitrile/ammonium acetate buffer as mobile phase for trimethoprim analysis. Considering that in a study carried out in China a maximum concentration of 5.2 ngL⁻¹ of trimethoprim was found in drinking water (Ben et al., 2020), it could be inferred that our method did not present adequate sensitivity for the determination of this compound.

Internationally, European countries are leading legislative actions concerning emerging contaminants. The European Commission has established a watch list of substances that may pose a risk to the aquatic environment but whose current information is limited to conclude its significance. Among the chemical compounds listed are two of those included in this study: sulfamethoxazole and trimethoprim (Implementing Decision (EU), 2022/1,307). Diclofenac was previously listed, in 2015 (OECD, 2019), and is currently regulated in Switzerland, with a maximum concentration of 50 ngL⁻¹ in surface waters (Swiss Federal Law, 2023).

In Ecuador, our results point towards several steps that must be taken to ensure water quality. One of the most important is the increase of wastewater treatment plants and the provision of various technologies to effectively remove or reduce polluting substances, including emerging contaminants. Wingfield et al. (2021) point out the importance of sewage treatment in ensuring water access in Ecuador, while also advocating for increased legislative support for ecosystem preservation. Along the same lines, Martínez-Moscoso et al. (2018) provide an overview of the current water laws landscape in Ecuador, emphasizing that water provision, particularly drinking water, shall be a public service. However, their study demonstrates that in practice, to comply with this precept, local governments pass on operating costs to final users, resulting in an increased cost to the population in rural areas. If wastewater treatment plants are installed in these municipalities, the cost of water services will rise, and if more advanced treatments are installed, the cost will rise even further. Hence, financial support from other sources is an important aspect to consider along with the cost of advanced treatment technologies. The Organization for Economic Cooperation and Development (2019) reports that from the currently available methods for pharmaceuticals removal from water, reverse osmosis and ozone have similar efficiency, but the latter has half of the cost per cubic meter.

The results from 179 samples collected in five Ecuadorian cities indicate that caffeine is one of the most prevalent emerging

compounds in drinking water. We have pointed out the growing need to monitor ECs in various bodies of water to anticipate and effectively mitigate the environmental risks that derive from them. Protection measures for drinking water sources should include strict regulations for houses, hotels, restaurants, and entertainment venues nearby. Of paramount importance are the implementation of adequate garbage disposal and wastewater treatment plants to avoid infiltration of liquids that endanger the quality of the source water. An update of the water treatment systems is also necessary, especially in the coastal cities that take river water as a source of water to be made drinkable. Future research should also consider extended monitoring campaigns and including drug metabolites in the analysis, which may appear due to reactivity in treatment plants, metabolism in the human body, or environmental degradation. These details will be critical for taking informed legislative action at the national level.

Data availability statement

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: https://osf.io/2mg87/?view_only=ea727d3ae65a40b0b81bad3cc1ee8327

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EJ-N: methodology, investigation, formal analysis, writing–original draft, writing–review and editing. IC-A: conceptualization, methodology, investigation, formal analysis, funding acquisition, supervision, writing–original draft, writing–review and editing. JM: methodology, investigation, formal analysis, validation, supervision, writing–original draft, writing–review and editing. VP-V: methodology, formal analysis, funding acquisition, writing–original draft, writing–review and editing. MC: methodology, validation, writing–review and editing. MC: methodology, formal analysis, writing–review and editing. EM-V: methodology, formal analysis, writing–review and editing. DL-A: methodology, investigation, writing–original draft. AP: investigation, writing–original draft. NM:

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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