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Spatial differences and temporal changes in illicit drug use in Europe quantified by wastewater analysis

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ABSTRACT

Aims To perform wastewater analyses to assess spatial differences and temporal changes of illicit drug use in a large European population. **Design** Analyses of raw wastewater over a 1-week period in 2012 and 2013. **Setting and Participants** Catchment areas of wastewater treatment plants (WWTPs) across Europe, as follows: 2012: 25 WWTPs in 11 countries (23 cities, total population 11.50 million); 2013: 47 WWTPs in 21 countries (42 cities, total population 24.74 million). **Measurements** Excretion products of five illicit drugs (cocaine, amphetamine, ecstasy, methamphetamine, cannabis) were quantified in wastewater samples using methods based on liquid chromatography coupled to mass spectrometry. **Findings** Spatial differences were assessed and confirmed to vary greatly across European metropolitan areas. In general, results were in agreement with traditional surveillance data, where available. While temporal changes were substantial in individual cities and years (P ranging from insignificant to $<10^{-3}$), overall means were relatively stable. The overall mean of methamphetamine was an exception (apparent decline in 2012), as it was influenced mainly by four cities. **Conclusions** Wastewater analysis performed across Europe provides complementary evidence on illicit drug consumption and generally concurs with traditional surveillance data. Wastewater analysis can measure total illicit drug use more quickly and regularly than is the current norm for national surveys, and creates estimates where such data does not exist.

Keywords Amphetamine, cannabis, cocaine, drugs of abuse, ecstasy, methamphetamine, sewage.

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INTRODUCTION

Illicit drug use is a covert and hidden activity that presents methodological challenges for drug surveillance

systems. Questionnaire-based survey methods have traditionally been an important component of the approaches employed to monitor drug use, but it is recognized that these methods are not sufficient to monitor

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trends in drug use adequately and quickly, and require complementary data from other sources [1,2]. The analysis of the excretion products of illicit drugs in wastewater [wastewater analysis (WWA)] has been explored since 2008 as an additional approach for estimating illicit drug use within specified regions, i.e. the catchment areas of wastewater treatment plants (WWTP) [3,4]. While the approach cannot provide information on the behaviour of single users and on their demographics, there are a number of ways in which WWA can complement other survey methods and provide additional information to understand the illicit drug situation more clearly. Wastewater data can be obtained within short time-frames, are not prone to response biases and can help in identifying the spectrum of illicit drugs being used by a population. This is potentially important, given the emergence of new psychoactive substances [5]. Drug users are often unaware of the actual substance or mix of substances they are consuming, which makes self-report data unreliable. Wastewater analysis is therefore a potential approach to detect and estimate the use of new psychoactive substances; however, it should be noted that more information is necessary regarding their biotransformation pathways.

Wastewater analysis can provide information on daily, weekly, monthly and annual variations in illicit drug use. The weekly profile of cocaine and amphetamine-like stimulants use has already been assessed by collecting consecutive daily wastewater samples, which revealed higher use of these substances during weekends [6–12]. The monitoring of temporal trends in illicit drug consumption over a longer period of time (months) by WWA has been evaluated in three studies, and the major conclusions were that there was typically an increase of illicit drug use during holiday periods [11,13,14]. Wastewater analysis was further applied to detect yearly trends in illicit drug consumption in Italy and Australia [15,16]. In conclusion, this approach can provide important and timely information on short- and long-term trends in illicit drug use.

Wastewater studies in different countries have also detected regional variations in illicit drug use [17–22]. The influence of urbanization on the use of illicit drugs was evaluated in Oregon (USA) and South Australia and Queensland (Australia), concluding that the use of illicit drugs was higher in urban regions compared to more rural areas [9,14,23]. Wastewater analysis has also been applied to detect transnational differences in illicit drug use. The consumption of five substances was evaluated by analysing wastewater from 19 European cities for a 1-week period in 2011 [24]. Wastewater analysis can thus complement survey methods for a clearer understanding of actual spatial differences and temporal changes in illicit drug use.

However, until now no international study has been performed covering multiple countries over multiple years with a common protocol and adequate quality control measures. Therefore, the aims of this study were to:

- 1 collect wastewater samples from multiple European locations in 2012 and 2013;
- 2 calculate population-normalized mass loads of benzoylecgonine [BE; as indicator for cocaine (COC) use], amphetamine (AMP), methamphetamine (METH), ecstasy [3,4-methylenedioxy-methamphetamine (MDMA)] and 11-nor-9-carboxy-delta9-tetrahydrocannabinol [THC-COOH; as indicator for tetrahydrocannabinol (THC) use]; and
- 3 perform analytical quality control through inter-laboratory tests.

METHODS

Sewer system characterization

Relevant information for each WWTP catchment was gathered systematically by means of a standardized questionnaire. An extended version of the questionnaire developed for earlier studies [24,25] was used (Supporting information, Appendix S1). It comprises more than 50 questions classified according to importance. The number of the most important questions per category is indicated in brackets (year 2012/year 2013): General information (1/1), Catchment and population (2/5), Sewer system (2/2), WWTP influent (1/1), Sampling (5/5), Flow meter (3/3), Sample handling (9/9), Monitoring period (5/5).

Sampling and analysis

A 1-week period was targeted in 2012 (17–23 April) and 2013 (6–12 March). Daily 24-hour composite raw wastewater samples were collected over 7 consecutive days. Considering stability, metabolism and unambiguous indication of drugs actually having been consumed, the most suitable target residues were targeted: BE, AMP, METH, MDMA and THC-COOH [4]. It should be noted that the consumption of COC and THC was monitored through the analysis of their main metabolite because of higher concentrations and higher stability in wastewater.

Samples were spiked with isotope-labelled internal standards, either filtered and extracted immediately on solid-phase extraction cartridges or frozen at -20°C until analysis. Each laboratory used fully validated analytical methods: target compounds present in the liquid phase of the wastewater were quantified in final extracts or with direct injection applying liquid chromatography coupled to tandem mass spectrometry or high-resolution mass spectrometry [25].

For quality assurance, each laboratory participated in yearly inter-laboratory tests (de Voogt *et al.*, unpublished). External quality control samples were evaluated (one standard in methanol and two fortified raw wastewater samples). A reliable estimation of the method limit of quantification (LOQ) was performed by evaluating the signal-to-noise ratio in these samples. In 2012, one of 14 laboratories did not meet the requirements for any compound in the inter-laboratory test and was excluded. In 2013, only METH results of one of 15 laboratories had to be excluded.

Calculations

Daily mass loads (g/day) of drug residues entering the WWTPs were calculated by multiplying measured concentrations (ng/L) in daily samples with the corresponding wastewater volumes (L/day). To compare cities of different sizes, mass loads are normalized by the population size of the catchment (mg/1000 people/day). The estimated consumption of COC (section Benzoylecgonine) was back-calculated from the population-normalized mass loads of BE using a correction factor of 3.59 that takes the urinary excretion rate of COC into account for different dosages and routes of administration [25].

Uncertainty assessment

Mainly four components of uncertainty may affect the estimation of population-normalized drug loads: sampling (U_S), chemical analysis (U_C), flow rate measurement (U_F) and population estimation (U_P). Because the focus of this study is on mass loads in wastewater, uncertainties related to excretion rates and biodegradation in sewers are not considered. When estimating the overall uncertainty U_T of a mean value over an n -day monitoring period, uncertainty components that are random and independent on every day will be reduced by \sqrt{n} . This applies to U_S , as each sample is collected physically independent of the day before. All other components cannot be reduced by \sqrt{n} : (i) population is only estimated once, (ii) chemical analysis is carried out for all samples in one batch, and (iii) if a flow meter measures flows systematically incorrectly, it will be in the same direction every day. All components can be considered as independent. As long as U_S , U_C and $U_F \leq 30\%$ and $U_P \leq 10\%$ [relative standard deviation (RSD)], an estimation of U_T is valid with an approximative formula (e.g. [26]). A Monte Carlo simulation was used to avoid underestimating U_T systematically because a conservative estimate of U_P in our study is 20% (see Supporting information, Appendix S2).

RESULTS

Table 1 lists participating cities: in 2012, 25 WWTPs in 11 countries were included (23 cities, total population

11.50 million); in 2013, there were 47 WWTPs in 21 countries (42 cities, total population 24.74 million). For comparison, 2011 data [24] were also used (21 WWTPs in 11 countries; 19 cities, total population 14.12 million). Figures 1–5 summarize all results. Countries are ordered based on average loads over all years. The numbers in brackets indicate cities' overall ranks. While absolute variability within 1-week periods (grey range) is obviously higher for high loads, relative variability is not substantially different throughout the entire load range and may vary from year to year, even within a location. The colour of the lines between the means indicate whether the change from 1 week in 1 year to 1 week in another year is significant (Wilcox, $\alpha = 0.05$). Table 2 summarizes overall means, separately for cities that participated in all 3 years (cities in bold type in Figs 1–5) and for all cities per year (excluding cities that exhibited explainable anomalies, i.e. cities in italic type in Figs 1–5). Concentration values that were <LOQ were treated as follows: (1) if all values at a location for a certain compound were <LOQ, loads were set to zero; (2) if at least one value was >LOQ, values <LOQ were replaced with $0.5 \times \text{LOQ}$. Dashed grey lines indicate a population-weighted overall mean for 2013 (all cities except cities in italics). When weekly patterns were evaluated in 2012, previous findings were confirmed, i.e. higher loads on weekends for BE, and MDMA and no substantial variation for AMP, METH and THC-COOH [24] (see Supporting information, Appendix S4).

Benzoylecgonine

The highest weekly mean BE loads in the period 2011–13 were observed in wastewater from Amsterdam, Antwerp, London and Zurich and were between 400–850 mg/1000 people/day (Fig. 1). Loads were also relatively high (between 200–550 mg/1000 people/day) in Barcelona, Basel, Geneva, Utrecht and Eindhoven. The lowest values (<100 mg/1000 people/day) were observed in locations from northern, eastern and southern Europe. These results suggest a clear geographical difference in COC consumption, with higher use in western Europe. This is further demonstrated when BE loads in locations from Germany are evaluated. Loads in Dresden (eastern Germany) are negligible, similar to the amounts seen in the Czech Republic, while loads in Dortmund (western Germany) are comparable to the loads observed in the Belgian, Dutch and Swiss cities.

The overall population-weighted mean loads of BE for the 16 locations included in all 3 years were almost identical (Table 2). This suggests a stable use of COC in the investigated locations in the period 2011–13. Location-specific results from 2011, 2012 and 2013 are generally in agreement (Fig. 1); however, in some cases, variations

Table 1 Summary of participating cities and wastewater treatment plants (WWTP). More detailed information can be found in Supporting information (Appendix S3), which includes raw data and answers from the questionnaire.

Country	City	WWTP	Population of the city under investigation [ⓐ]	① Estimated population in WWTP catchment				Targeted 1-week monitoring period [✓] (n = 7 days)				Loss of wastewater [ⓑ]	Com-miters [ⓒ]	Special events [ⓓ]
				2011	2012	2013	method (year)	2011	2012	2013	Mar 6–12			
BA	Sarajevo	Butle	C: 291 422; M: 515 012 (W)			130 000	b (2013)		Mar 9–15	Apr 17–23	Mar 6–12	✓	ⓓ	R (2013)
BE	Antwerp D.	Deurne	498 473 (E 2011)	213 876	a (2011)	213 876	a (2012)		✓	✓	✓	✓	ⓓ	
	Antwerp Z.	Zuid	498 473 (E 2011)	117 200	a (2010)	130 218	a (2011)		✓	✓	✓	✓	ⓓ	
	Brussels	Noord	1136 778 (E 2011)	1 027 300	b (2011)	953 987	b (2012)		✓	✓	✓	✓	ⓓ	
	Geraardsbergen	Geraardsbergen	32 629 (W)			29 047	c (2011)		March 8–13				ⓓ	
	Koksijde	Wulpen	31 207 (W)			78 441	a (2012)		March 21–27				ⓓ	
CH	Ninove	Ninove	37 295 (W)			36 179	c (2013)		✓	✓	✓	✓	ⓓ	
	Basel	ProRheno	C: 195 743 (L 2013)	260 000	c (2012)	260 000	c (2012)		✓	✓	✓	✓	ⓓ	R (2012)
	Berne	Region Bern	C: 137 818 (L 2012)	206 655	c (2012)	206 700	c (2012)		April 21–27	April 22–28	✓	✓	ⓓ	R (2012)
	Geneva	SIG	C: 194 458 (L 2013)	410 486	c (2012)	410 500	c (2012)		✓ (n = 5)	✓	✓	✓	ⓓ	R (2012)
	St Gallen	Au and Hofen	C: 74 070 (L 2013)	89 000	c (2012)	89 000	c (2012)		✓	✓	✓	✓	ⓓ	R (2012)
CY	Zurich	Werdhölzli	C: 394 012 (L 2012)	410 000	c (2012)	410 000	c (2012)		✓	✓	✓	✓	ⓓ	Y (2012)
	Nicosia	Pano Delfera	C: 234 200 (E 2009)	28 000	c (2012)	28 000	c (2012)		March 21–27	March 21–27	✓	✓	ⓓ	Y (2013)
	Limassol	Amathus	185 100 (E 2009)	272 000	c (2010)	272 000	c (2010)		March 21–27	March 21–27	✓	✓	ⓓ	Y (2013)
	Budweis	COV	93 620 (E 2011)	112 000	d (2010)	110 300	d (2013)		✓	✓	✓	✓	ⓓ	Y (2011)
	Prague	UCOV	1241 664 (E 2011)	1 300 000	c (2011)	1 300 000	c (2011)		✓	✓	✓	✓	ⓓ	Y (2011)
DE	Dortmund	Deusen	580 956 (E 2012)	371 788	c (2010)	371 788	c (2010)		March 13–19	March 13–19	✓	✓	ⓓ	
	Dülmen	Dülmen	46 071 (W)	34 495	c (2010)	34 495	c (2010)		✓	✓	✓	✓	ⓓ	
	Dresden	Kaditz	529 781 (E 2012)	593 050	c (2012)	593 050	c (2012)		✓	✓	✓	✓	ⓓ	
DK	Copenhagen	Lynetten	501 285 (E 2003)	531 000	c (2009)	531 000	c (2009)		March 16–22	March 16–22	✓	✓	ⓓ	Y (2012/13)
	Barcelona	Baix Llobregat	C: 1 620 943; M: 3 202 571 (E 2012)	1 162 000	c (2007)	1 162 000	c (2010)		March 6–13 (n = 7)	March 6–13 (n = 7)	✓	✓	ⓓ	R (2013)
ES	Castellon	Castellon de la Plana	C: 180 204 (E 2012)	170 600	a (2010)	204 878	b (2012)		✓	✓	February 20–26	✓	ⓓ	
	Santiago	Silvota	95 671 (E 2012)	136 500	d (2010)	136 500	d (2010)		✓ (n = 6)	✓	✓	✓	ⓓ	
	Valencia	Pinedo I + II [ⓐ] and QB [ⓐ]	C: 797 028 (E 2012); M: 1 353 250 (L 2013)	1 839 000	a [ⓐ] /b [ⓐ] (2011)	1 357 952	e [ⓐ] (2011)/b [ⓐ] (2013)		✓	✓	✓	✓	ⓓ	R (2011/13)

Table 1 Cont.

Country	City	WWTP	Population of the city under investigation [ⓐ]	Estimated population in WWTP catchment				Targeted 1-week monitoring period [✓] (n = 7 days)				Loss of wastewater [ⓑ]	Com-muters [ⓓ]	Special events [ⓔ]
				2011	method (year)	2012	method (year)	2013	method (year)	2011 Mar 9–15	2012 Apr 17–23			
FI	Helsinki		M: 1 022 139 (E 2009)	780 000 a (2009)	780 000 a (2009)	780 000 a (2009)	✓	✓	✓	△	-	-		
	Vikimäki						✓ (n = 6)	✓	✓	△	-	-	R (2012)	
	Kakola		178 630 (E 2012)	*275 000 d (2011)	275 000 d (2011)	275 000 d (2011)	✓	✓	✓	△	-	-	Y (2012)	
FR	Seine Centre*/Grésillon**		C: 2 243 718 (E 2010), M: 6 507 783 (E 2006)	+774 600 b (2011)	+245 500 f (2012)	*1 004 000 b (2013)	✓	✓	✓	●	▷			
	Beckton		8 174 000 (L 2011)	3 400 000 a (2010)	3 400 000 e (2010)	3 400 000 e (2010)	✓ (n = 6)	✓	✓	-	-	-		
GR	Athens		M: 2 989 023 (E 2009)	650 000 c (2001)	650 000 c (2011)	650 000 c (2011)	✓	✓	✓	△	-	-		
HR	Zagreb		C: 688 163, M: 1 107 623 (W)	650 000 c (2001)	650 000 c (2011)	650 000 c (2011)	✓	✓	✓	-	-	-		
IT	Milano		1 295 705 (E 2009)	1 250 000 c (2010)	1 100 000 b (2012)	1 149 477 b (2013)	✓	✓	✓	△	▷			
	Nosodo		779 808 (E 2011)	694 800 b (2011)	769 000 c (2010)	769 000 c (2010)	✓	✓	✓	△	▷		Y (2012)	
NL	Amsterdam		1 250 000 c (2010)	1 250 000 c (2010)	1 100 000 b (2012)	1 149 477 b (2013)	✓	✓	✓	△	▷			
	West		779 808 (E 2011)	694 800 b (2011)	769 000 c (2010)	769 000 c (2010)	✓	✓	✓	△	▷		Y (2012)	
Eindhoven			216 036 (E 2011)	448 700 b (2011)	450 300 c (2005)	450 300 c (2005)	✓	✓	✓	-	-	-	Y (2011/12)	
	Utrecht		311 367 (E 2011)	297 000 b (2011)	300 000 c (2010)	300 000 c (2010)	✓ (n = 5)	✓	✓	●	-	-	Y (2011)	
NO	Oslo		599 230 (E 2011)	557 000 c (2009)	557 000 c (2009)	576 000 c (2012)	✓	✓	✓	△	▷			
	VEAS						✓ (n = 6)	✓	✓	△	▷			
PT	Lisbon		C: 537 412, M: 1 860 256 (E 2012)	557 000 c (2009)	557 000 c (2009)	426 964 c (2011)	✓	✓	✓	●	▷			
	Alcantara						✓ (n = 6)	✓	✓	●	▷			
RO	Cluj Napoca		304 802 (E 2011)	315 000 c (2009)	664 441 c (2011)	664 441 c (2011)	✓	✓	✓	△	-	-		
	Cluj Napoca						✓	✓	✓	-	-	-		
RS	Belgrade Sewer		C: 1 232 731, M: 1 659 440 (W)	315 000 c (2009)	664 441 c (2011)	664 441 c (2011)	✓	✓	✓	△	-	-		
	Outlet Danube						✓	✓	✓	△	-	-		
SE	Novi Sad Sewer		M: 341 625 (W)	315 000 c (2009)	664 441 c (2011)	664 441 c (2011)	✓	✓	✓	△	-	-		
	Outlet Danube						✓	✓	✓	△	-	-	R (2013)	
Stockholm	Ryaverket		513 751 (E 2010)	315 000 c (2009)	664 441 c (2011)	664 441 c (2011)	✓	✓	✓	△	-	-		
	Henriksdals (only 1 of 2 inlets)		C: 847 073, M: 1 550 208 (E 2010)	315 000 c (2009)	664 441 c (2011)	664 441 c (2011)	✓	✓	✓	△	-	-		
SK	Umeå		115 473 (E 2010)	115 800 c (2010)	115 800 c (2010)	115 800 c (2010)	✓	✓	✓	△	▷			
	Bratislava		415 589 (L 2012)	115 800 c (2010)	115 800 c (2010)	440 000 c (2011)	✓	✓	✓	●	▷		Y (2013)	
Piestany	Petržalka (2 WWTPs)		29 660 (W)	30 000 c (2011)	30 000 c (2011)	30 000 c (2011)	✓	✓	✓	●	▷		Y (2013)	
	Piestany						✓	✓	✓	●	▷		Y (2013)	

ⓐ Population of entire city/region. C: city; M: metropolitan, greater region. (E: W, L): Eurostat, Wikipedia and local bureau for population statistics (year). ⓑ Method for population estimation in WWTP catchment (year of estimate). a. Influent nutrient load over corresponding calendar year; b. influent nutrient load over actual sampling period; c. census; d. house connections/drinking water subscribers; e. values adopted from previous estimation; f. WWTP different from 2011/13 but wastewater from same catchment (central collection with subsequent distribution to different WWTPs). *Population estimate indicated in [24] was erroneous and population-normalized consumption estimates are corrected with updated value. ⓓ Loss of wastewater (exfiltration, questionnaire 2013). △: no loss expected; ●: loss indicated (unknown amount or <20%); ●: loss >20% expected; -: information missing. ⓔ Commuters (work days versus weekend, questionnaire 2013). ▷: No substantial net population increase/decrease due to commuters; ◻: net increase of population on workdays; -: information missing. ⓕ Special events during/adjacent to monitoring period. Y: please see Supporting information, Appendix S3 for type of event (year provided in brackets); R: rain before/during monitoring period (higher flows but no substantial effect on drug loads expected).

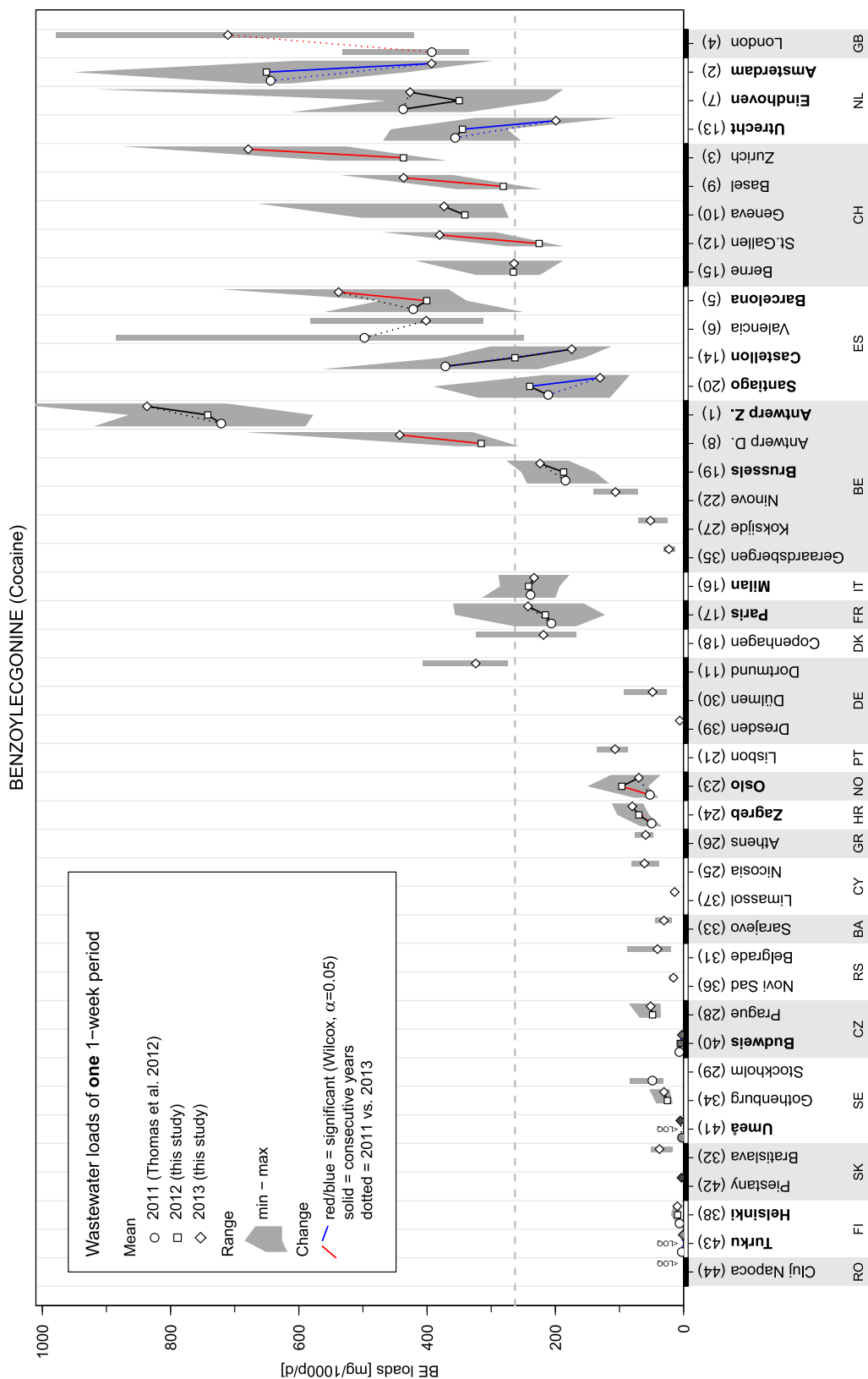


Figure 1 Population-normalized benzoylcegonine (BE) loads of a single 1-week period per year. See Table 1 for more information. <LOQ> concentrations in all daily samples were below limit of quantification (LOQ). **Grey dashed line:** 2013 overall mean of all participating cities. **Dot colour:** white: concentrations in all samples were above LOQ; grey shading: one or more concentrations were below LOQ and set to 0.5*LOQ (the darker the grey, the more concentrations were below LOQ). **Numbers in brackets:** cities' rank (average over all available years). Cities in **bold type** participated in all 3 years and were used to calculate annual overall means (see Table 2). All P-values can be found in Supporting information, Appendix S3

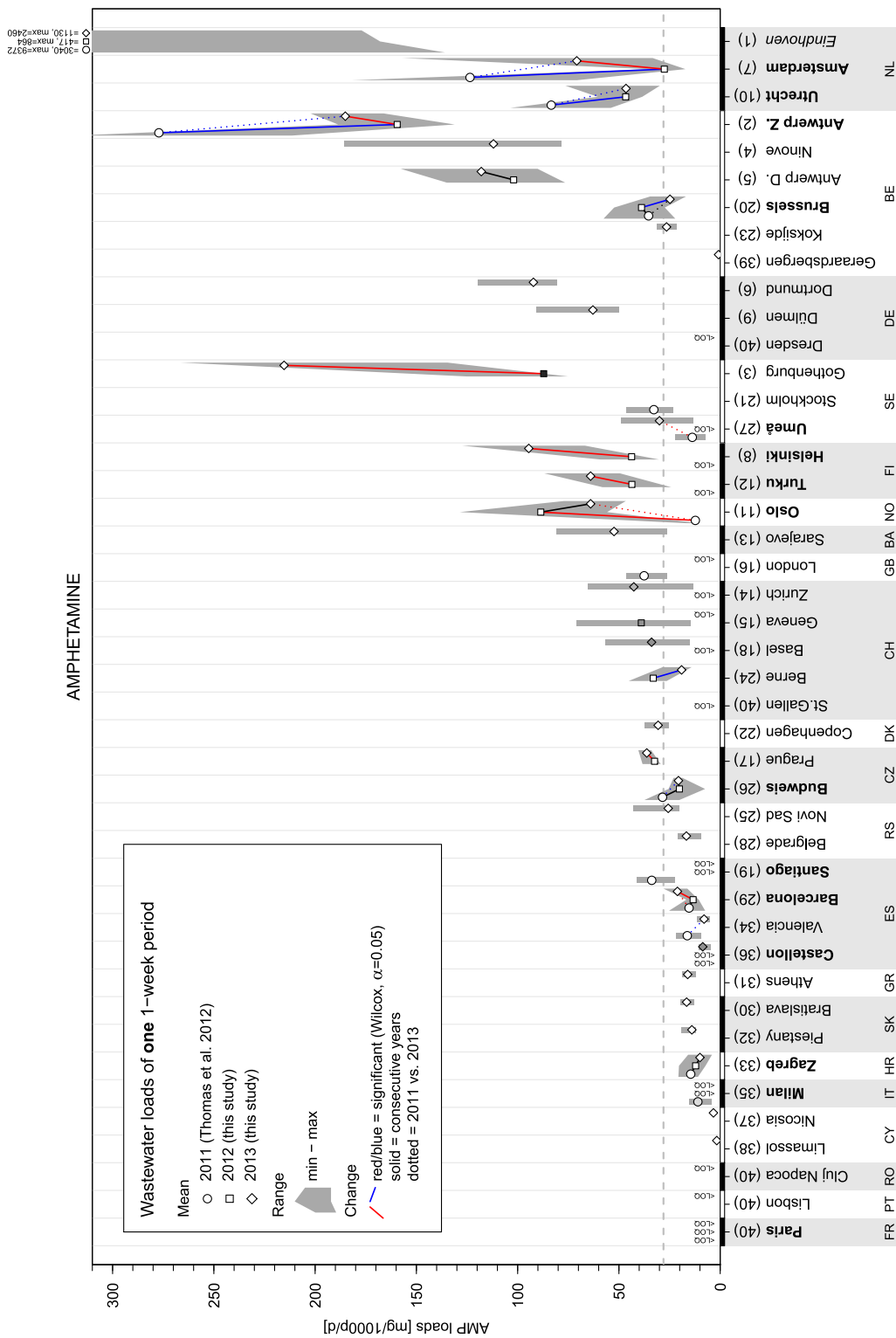


Figure 2 Population-normalized amphetamine (AMP) loads of a single 1-week period per year. See Table 1 for more information. <LOQ concentrations in all daily samples were below limit of quantification (LOQ). **Grey dashed line:** 2013 overall mean of all participating cities (except Eindhoven). **Dot colour:** white: concentrations in all samples were above LOQ; grey shading: one or more concentrations were below LOQ and set to 0.5%LOQ (the darker the grey, the more concentrations were below LOQ). **Numbers in brackets:** cities' rank (average over all available years). Cities in **bold type** participated in all 3 years and were used to calculate annual overall means (see Table 2). Cities in **italic type** exhibited abnormal high values in at least 1 year (see text for more details). All P-values can be found in Supporting information, Appendix S3

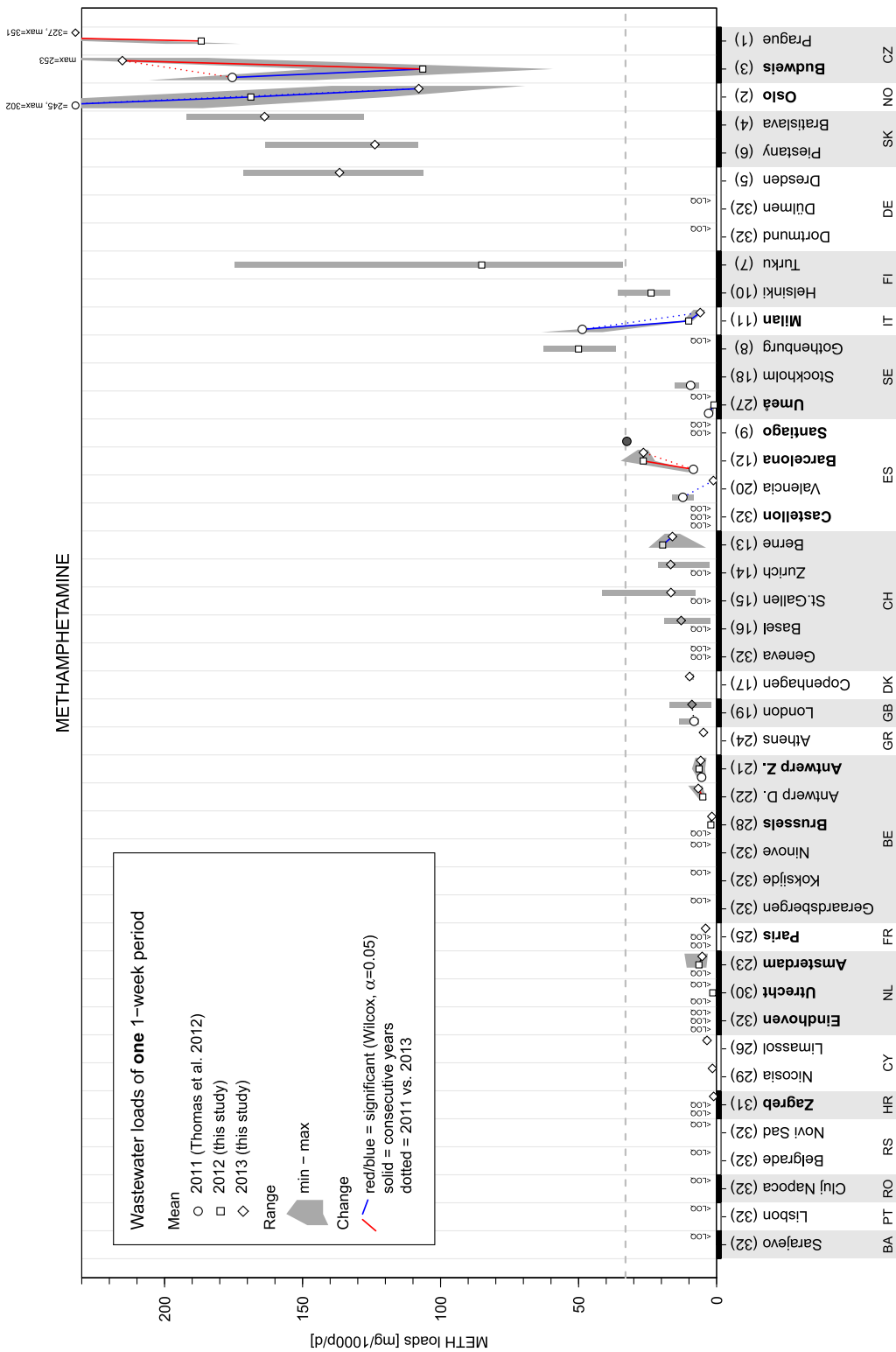


Figure 3 Population-normalized methamphetamine (METH) loads of a single 1-week period per year. See Table 1 for more information. <LOQ: concentrations in all daily samples were below limit of quantification (LOQ). **Grey dashed line:** 2013 overall mean of all participating cities. **Dot colour:** white: concentrations in all samples were above LOQ; grey shading: one or more concentrations were below LOQ and set to 0.5*LOQ (the darker the grey, the more concentrations were below LOQ). **Numbers in brackets:** cities' rank (average over all available years). **Cities in bold type** participated in all 3 years and were used to calculate annual overall means (see Table 2). All P-values can be found in Supporting information, Appendix S3

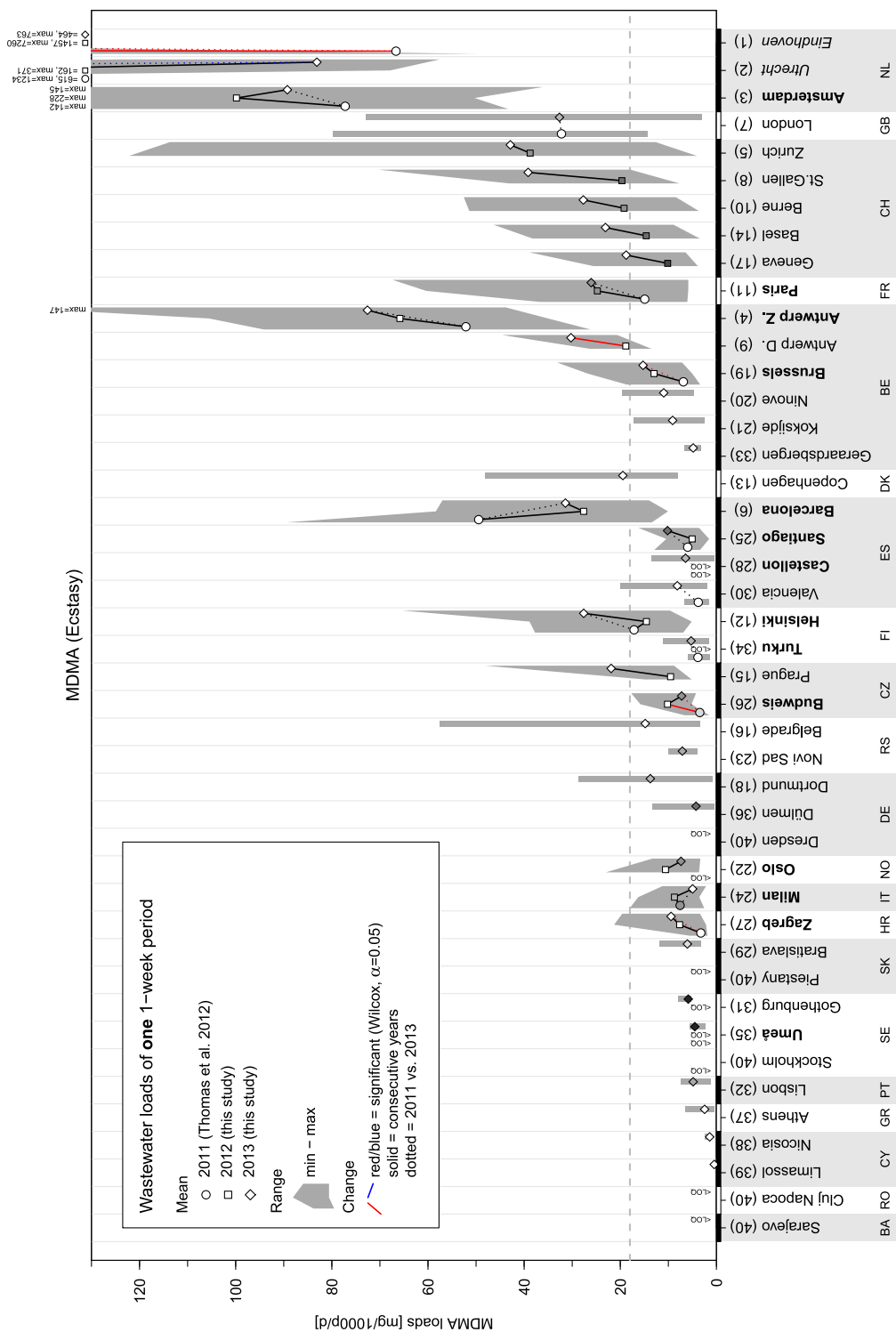


Figure 4 Population-normalized 3,4-methylenedioxy-methamphetamine (MDMA) loads of a single 1-week period per year. See Table 1 for more information. <LOQ: concentrations in all daily samples were below limit of quantification (LOQ). **Grey dashed line:** 2013 overall mean of all participating cities (except Utrecht and Eindhoven). **Dot colour:** white: concentrations in all samples were above LOQ; grey shading: one or more concentrations were below LOQ and set to 0.5*LOQ (the darker the grey the more concentrations were below LOQ). **Numbers in brackets:** cities' rank (average over all available years). Cities in **bold type** participated in all 3 years and were used to calculate annual overall means (see Table 2). Cities in **italic type** exhibited abnormal high values in at least 1 year (see text for more details). All P-values can be found in Supporting information, Appendix S3

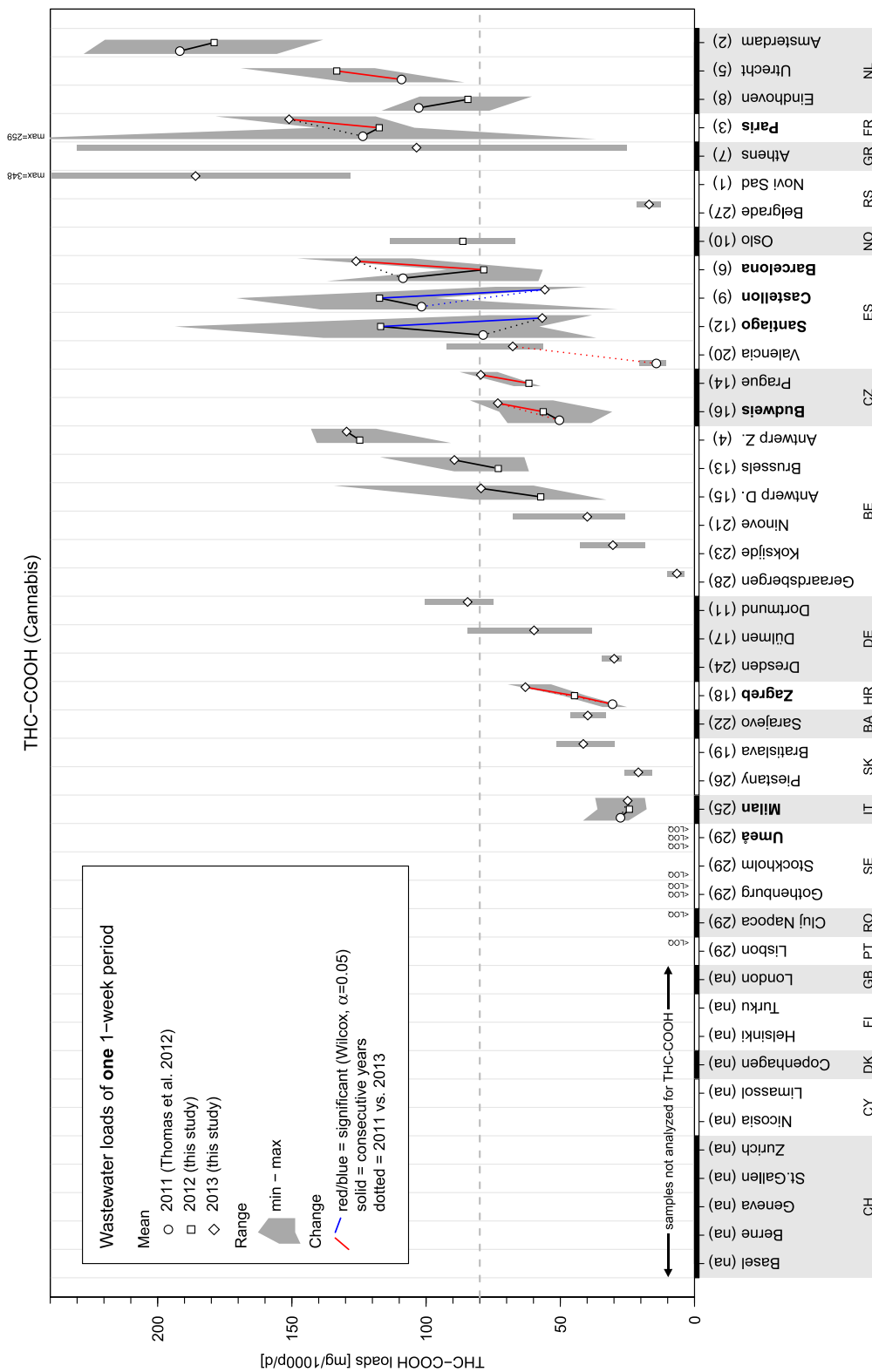


Figure 5 Population-normalized 11-nor-9-carboxy-delta9-tetrahydrocannabinol (THC-COOH) loads of a single 1-week period per year. See Table 1 for more information. $\lt; LOQ$ concentrations in all daily samples were below limit of quantification (LOQ). **Grey dashed line:** 2013 overall mean of all participating cities. **Dot colour:** white: concentrations in all samples were above LOQ; Grey shading: one or more concentrations were below LOQ and set to 0.5*LOQ (the darker the grey, the more concentrations were below LOQ). **Numbers in brackets:** cities' rank (average over all available years). Cities in **bold type** participated in all 3 years and were used to calculate annual overall means (see Table 2). All P-values can be found in Supporting information, Appendix S3

Table 2 Population-weighted overall mean loads (units = mg/1000p/d). The loads in cities with all concentration values <LOQ were set to 0. Loads range from (close to) 0 up to several 10–100 mg/1000 person/day among cities, which implies large standard deviation (SD) or 95% confidence interval (CI) for all substances' overall means. Therefore, significance of changes cannot be meaningfully assessed for overall means and is assessed at cities' individual levels only (see Figs 1–5 and Supporting information, Appendix S3)

	BE		MDMA		AMPH		METH		THC-COOH	
	a	b	a	b	a	b	a	b	a	b
2011 {14,12}	249 [8.57]	311 [14.12]	21 [7.82]	21 [13.38]	29 [8.12]	30 [13.67]	31 [7.51]	22 [13.07]	71 [4.37]	69 [7.97]
2012 {11,50}	254 [7.94]	229 [11.50]	24 [7.19]	20 [10.75]	29 [7.49]	32 [11.05]	23 [6.89]	42 [11.50]	60 [3.73]	73 [9.07]
2013 {24,74}	247 [8.77]	263 [24.74]	25 [8.02]	18 [23.99]	34 [8.32]	28 [24.20]	17 [7.71]	33 [23.68]	87 [4.53]	77 [15.98]

^aOnly cities participating in all 3 years are considered. These cities are labelled in bold type in the corresponding figures 1–5. Cities with 'explainable anomalies' for a particular substance are excluded from the calculation of overall means and labelled in italic type (even if the anomaly occurred only in 1 year). ^bAll cities participating in the corresponding year are considered except the ones that were already excluded due to 'explainable anomalies' in option a. Cities with 'explainable anomalies' for a particular substance are excluded from the calculation of overall means and labelled in italic type (even if the anomaly occurred only in 1 year). {} Total population in millions monitored (please note: not all substances were measured in all cities). [] Population in millions contributing to the corresponding overall mean. BE = benzoyllecgonine; MDMA = 3,4-methylenedioxy-methamphetamine; AMPH = amphetamine; METH = methamphetamine; THC-COOH = 11-not-9-carboxy-delta9-tetrahydrocannabinol.

occurred. An increase in BE loads from 2012 to 2013 was observed in the Belgian and Swiss locations, while a decrease was observed in two Dutch locations (Utrecht and Amsterdam).

Besides the high variation of mean BE loads observed across Europe, this study also highlights differences among locations within countries. Results from Belgium, Czech Republic, Germany, Serbia, Slovakia, Sweden and Switzerland suggest that the consumption of COC is lower in smaller towns compared to larger cities (Table 1, Fig. 1). Qualitatively, this is in agreement with studies investigating more locations within a country [17–22], although some of these rely on grab samples or single days only. The difference between Dresden and Dortmund, two cities of similar size, is attributable to their geographic location within Germany, as discussed previously.

The population-weighted mean COC consumption, calculated from BE loads (see Calculations), for locations included in all study years is similar between years and varies from 887 mg/1000 people/day in 2013 to 912 mg/1000 people/day in 2012. With 366 million people living in the urbanized regions of the European Union and a mean purity of 39% [standard deviation (SD) = 12%] [27,28], a rough extrapolation would imply that 832 kg of street purity COC per day is consumed by the urbanized population in the European Union in 2013.

Amphetamine and methamphetamine

Because AMP is a urinary metabolite of METH and as AMP in wastewater could subsequently result from the use of METH, loads of both substances in wastewater have to be evaluated in parallel. Moreover, the use of certain prescription drugs, such as selegiline, may also result in traces of AMP and METH in wastewater following its metabolism; however, prescription rates indicate that any contribution would typically be <1% of the total AMP signal [24,29]. The most frequent amphetamine-like substance detected in the majority of the investigated locations was AMP. The highest AMP loads were found in Belgium and the Netherlands, followed by locations in northern Europe and western Germany. The locations with the highest METH loads were found in the Czech Republic, Slovakia, eastern Germany and northern Europe, while the observed METH loads in the rest of the studied locations was low to even negligible (Figs 2 and 3). The presented results suggest an apparent geographical difference in the use of the amphetamine-like stimulants. The consumption of AMP is more widespread in western Europe, while the use of METH is clearly shown in northern Europe, Slovakia and Czech Republic. The German results confirm the aforementioned trend in the

use of amphetamine-like substances. In Dülmen and Dortmund (West), relatively high AMP and negligible METH use was observed, while for Dresden (East, proximity to Czech Republic) the opposite was found.

The weighted mean of METH loads for the cities that were included in all study years declined by 45% from 2011 to 2013 (Table 2), due to some location-specific changes. For AMP, the weighted mean of the cities included in the 3 years is similar (Table 2). In contrast to BE loads, the difference in AMP and METH loads between smaller towns and bigger cities within a country is less clear.

MDMA

The highest loads of MDMA were found in western European locations, while locations in northern, eastern and southern Europe presented substantially lower MDMA loads (Fig. 4). This pattern is comparable to BE and AMP, as demonstrated by the locations within Germany, with low MDMA loads in Dresden and higher loads in Dortmund.

The weighted mean of MDMA loads for the cities included in all 3 study years was stable (Table 2). No substantial changes in per capita MDMA loads between years for the individual locations were observed, with some exceptions (Fig. 4). The mass loads of MDMA from Eindhoven in 2012 and 2013 were much higher compared to 2011, and in Utrecht significantly higher loads for MDMA were observed in 2011 compared to 2012 and 2013. An explanation for these high loads in Utrecht (2011) and Eindhoven (2012) is most probably a release of unconsumed MDMA into the sewer system that was confirmed by specific enantiomeric profiling of the wastewater [30]. These outliers were not taken into account when assessing temporal changes. MDMA loads are generally higher in larger cities compared to smaller towns, as can be seen in different locations within Belgium, Finland, Germany, Serbia and Slovakia. A notable exception is St Gallen in Switzerland, which showed MDMA loads comparable to the larger city of Zurich.

THC-COOH

The determination of THC-COOH in wastewater poses some (pre-)analytical challenges, and as a result not all laboratories could report results for this THC metabolite. Furthermore, results from the performed inter-laboratory exercises revealed that participating laboratories that reported results for THC-COOH have comparable analytical methods (*Z*-scores within the limits), but because of some unknown pre-analytical losses, underestimations of the absolute amounts are probably made (de Voogt *et al.*, unpublished). In the present study, however, this is not a

real issue, because the focus lies on the relative comparison of THC-COOH loads.

In contrast to the other investigated substances, no clear geographical pattern could be observed for THC-COOH loads in the different European locations (Fig. 5). The values for Amsterdam were (expectedly) the highest, as Amsterdam is known for its coffee shops and because the Netherlands produces large amounts of herbal cannabis with a relatively high content of THC [31]. Also notable are the high loads observed in the city of Novi Sad, Serbia.

The weighted mean of THC-COOH loads for cities that were included in all 3 years showed some subtle variation, pointing out a variable cannabis use (amount or potency) between 2011 and 2013 (Table 2). No clear difference in THC-COOH loads between smaller towns and larger cities could be observed from the gathered data.

DISCUSSION

Comparison of wastewater results with surveillance data

Europe has an established multi-indicator system for drug surveillance that is based on standardized demand and supply information, as well as research and intelligence sources [32]. Prevalence estimates are derived from a mixture of survey results and indirect statistical methods that try to estimate the unobserved cases from registers of observed drug users, such as treatment attendees or arrestees [33]. These methods can provide information on the main classes of users, the frequency and mode of use of a drug as well as on the purity of the substances available on the market, while WWA can provide objective and timely information on the total amount of a drug used in a specific area. These methods are highly complementary and, if used together, can substantially improve the quality of information on drug use patterns.

In terms of prevalence at the population level, the findings from WWA are broadly in agreement, with respect to relative drug use levels, with existing estimates, although they are not directly comparable. The wastewater data, however, highlight the need to consider the contribution of high and low prevalence areas in the estimates of total drug use within a population. Due to differences in demographics, the ranking of the city-based estimates reported in this study do not necessarily have to agree with national survey-based estimates. This points to the need to collect contextual information for a meaningful interpretation of wastewater data. Future monitoring campaigns should therefore (i) include more cities with different demographics within a country and (ii) evaluate monitoring design strategies to find an

optimum among feasible logistics, sufficient quality control and representativeness for an entire year [34].

The spatiotemporal data on drug use data reported are largely, but not totally, in line with what is observed from surveys and other sources. The stable levels of COC suggested by the presented wastewater data differs from other demand and supply data, which report a decline in COC use [35]. With WWA, it is currently not possible to differentiate between smaller number of people using larger amounts or vice versa, or even evaluating differences in consumption due to changes in purity. The analysis on METH and AMP accords with other data sources. The use of METH is long established in the Czech Republic, Slovakia and eastern Germany [36], and more recently supply-side data point to an increased use of METH elsewhere, especially in Scandinavian countries where it has, at times, displaced AMP. The situation appears quite dynamic and largely supply-side-driven. The wastewater data reported here accords with, and complements, the existing analysis of this situation.

For both MDMA and cannabis use, the picture is less clear. High levels of MDMA and THC-COOH might be expected in the Dutch cities sampled, but it is surprising that MDMA stands out so prominently with respect to some of the other European cities. The most recent supply-side data suggest that there is more MDMA available on the European market, and it is interesting to note that there is no evidence of this from the wastewater data reported here. The findings for THC-COOH in Amsterdam are not too surprising, as it is known for its large non-resident population using cannabis.

Uncertainty assessment

Details on estimating U_S can be found in [37,38]. Applying the same scenario as in [25]—i.e. 1% of users in the population with two relevant, substance-related toilet flushes—results in a maximum of 20% for a daily value of U_S . An objective assessment of U_C was derived from inter-laboratory tests and does not exceed 30% (de Voogt *et al.*, unpublished). Operational accuracy of flow meters (U_F) still proves to be a challenge, and in this study was assumed conservatively to be 20% [39]. Despite advances in estimating U_P [40] it remains difficult to obtain a site-specific estimate, and in our study we assume 20% (RSD) as an average [25,40]. A conservative estimate of overall uncertainty for a 7-day average based on WWA is approximately 46% (RSD) for all substances and locations (see Supporting information, Appendix S2 for more details). A sensitivity analysis reveals that reducing all four uncertainty components U_i by approximately one-quarter ($U_S \approx U_F \approx U_P \approx 15\%$, $U_C \approx 23\%$) has the same effect as trying to eliminate only one U_i (e.g. $U_C \approx 0\%$); in both cases the overall uncertainty would be around 33%.

In areas with leaky sewers the results from WWA may tend towards an underestimation of actual illicit drug loads. A certain fraction of the wastewater and illicit drugs discharged from households may not arrive at the WWTP. Information on the potential amount of exfiltration can be found in Table 1. Furthermore, in cases where population size is estimated from nutrient loads in the wastewater stream, the population could be overestimated if industrial contributions are not properly subtracted. This would lead to an underestimation of population-normalized drug loads. In contrast, WWA results may tend towards an overestimation of population-normalized drug loads if the residential population only was used for normalization, but a net increase on workdays is effective due to commuters. This and additional information is provided in Table 1 and Supporting information, Appendix S3 for further data interpretation.

CONCLUSIONS

By successfully increasing the number of participating cities to 42 in 2013 (2011: 19, 2012: 23), this is now the biggest application of WWA covering 24.74 million people. The wastewater from approximately 8 million people was analysed for BE, AMP, METH and MDMA during a 1-week period over 3 consecutive years (approximately 4 million for THC-COOH). As such, this study provides the most actual evidence for the quantification of spatial differences and temporal changes in the consumption of illicit drugs across European regions. Relatively stable loads for all investigated substances were observed, except for METH (apparent decline in 2012). In general, spatial differences were in agreement with surveillance data, where available. Wastewater analysis provides the possibility to collect, and report, measurements more quickly and regularly than is the current norm for national surveys. Wastewater analysis provides a unique opportunity to obtain near-real-time data on illicit drug use and for future comparison with other surveillance data, or particularly where such data are missing. Therefore, it should be considered for implementation on an annual or even more frequent basis. Systematically gathering information on catchment characteristics (sewer system and population) seems as indispensable as inter-laboratory tests for a meaningful comparison of wastewater data, which requires concerted efforts of numerous partners and disciplines.

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Declaration of interests

None.

References

1. European Monitoring Centre for Drugs and Drug Addiction. Assessing Illicit Drugs in Sewage—Potential and Limitations of A New Monitoring Approach. Luxembourg: Office for Official Publications of the European Communities; 2008.

2. Griffiths P., Mounteney J. Drug trend monitoring. In: Miller P. G., Strang J., Miller P. M., editors. *Addiction Research Methods*. Oxford, UK: Wiley-Blackwell; 2010, pp. 337–54.
3. Zuccato E., Chiabrando C., Castiglioni S., Bagnati R., Fanelli R. Estimating community drug abuse by wastewater analysis. *Environ Health Perspect* 2008; **116**: 1027–32.
4. van Nuijs A. L. N., Castiglioni S., Tarcomnicu I., Postigo C., de Alda M. L., Neels H. *et al.* Illicit drug consumption estimations derived from wastewater analysis: a critical review. *Sci Total Environ* 2011; **409**: 3564–77.
5. United Nations Office on Drugs and Crime (UNODC). The challenge of new psychoactive substances. UNODC, Vienna, 2013.
6. Karolak S., Nefau T., Bailly E., Solgadi A., Levi Y. Estimation of illicit consumption by wastewater analysis in Paris area. *Forensic Sci Int* 2010; **200**: 153–60.
7. Metcalfe C., Tindale K., Li H., Rodayan A., Yargeau V. Illicit drugs in Canadian municipal wastewater and estimates of community drug use. *Environ Pollut* 2010; **158**: 3179–85.
8. Terzic S., Senta I., Ahel M. Illicit drugs in wastewater of the city of Zagreb (Croatia)—estimation of drug abuse in a transition country. *Environ Pollut* 2010; **158**: 2686–93.
9. Irvine R. J., Kostakis C., Felgate P. D., Jaehne E. J., Chen C., White J. M. Population drug use in Australia: a sewage analysis. *Forensic Sci Int* 2011; **210**: 69–73.
10. Reid M. J., Langford K. H., Morland J., Thomas K. V. Quantitative assessment of time dependent drug-use trends by the analysis of drugs and related metabolites in raw sewage. *Drug Alcohol Depend* 2011; **119**: 179–86.
11. van Nuijs A. L. N., Mougel J.-F., Tarcomnicu I., Bervoets L., Blust R., Jorens P. G. *et al.* Sewage epidemiology—a real-time approach to estimate the consumption of illicit drugs in Brussels, Belgium. *Environ Int* 2011; **37**: 612–21.
12. Bijlsma L., Emke E., Hernandez E., de Voogt P. Investigation of drugs of abuse and relevant metabolites in Dutch sewage water by liquid chromatography coupled to high resolution mass spectrometry. *Chemosphere* 2012; **89**: 1399–406.
13. Harman C., Reid M., Thomas K. V. *In situ* calibration of a passive sampling device for selected illicit drugs and their metabolites in wastewater, and subsequent year-long assessment of community drug usage. *Environ Sci Technol* 2011; **45**: 5676–82.
14. Lai F. Y., Bruno R., Hall W., Gartner C., Ort C., Kirkbride P. *et al.* Profiles of illicit drug use during annual key holiday and control periods in Australia: wastewater analysis in an urban, a semi-rural and a vacation area. *Addiction* 2013; **108**: 556–65.
15. Zuccato E., Castiglioni S., Tettamanti M., Olandese R., Bagnati R., Melis M. *et al.* Changes in illicit drug consumption patterns in 2009 detected by wastewater analysis. *Drug Alcohol Depend* 2011; **118**: 464–9.
16. Chen C., Kostakis C., Harpas P., Felgate P. D., Irvine R. J., White J. M. Marked decline in 3,4-methylenedioxymethamphetamine (MDMA) based on wastewater analysis. *J Stud Alcohol Drugs* 2011; **72**: 737–40.
17. Huerta-Fontela M., Galceran M. T., Martin-Alonso J., Ventura F. Occurrence of psychoactive stimulatory drugs in wastewaters in north-eastern Spain. *Sci Total Environ* 2008; **397**: 31–40.
18. van Nuijs A. L. N., Pecceu B., Theunis L., Dubois N., Charlier C., Jorens P. G. *et al.* Can cocaine use be evaluated through analysis of wastewater? A nation-wide approach conducted in Belgium. *Addiction* 2009; **104**: 734–41.

19. Kankaanpää A., Ariniemi K., Heinonen M., Kuoppasalmi K., Gunnar T. Use of illicit stimulant drugs in Finland: a wastewater study in ten major cities. *Sci Total Environ*; in press; 2014; doi:10.1016/j.scitotenv.2013.11.095
20. Vuori E., Happonen M., Gergov M., Nenonen T., Järvinen A., Ketola R. A. *et al.* Wastewater analysis reveals regional variability in exposure to abused drugs and opioids in Finland. *Sci Total Environ*; in press; 2014; doi:10.1016/j.scitotenv.2013.11.010
21. Östman M., Fick J., Näsström E., Lindberg R. H. A snapshot of illicit drug use in Sweden acquired through sewage water analysis. *Sci Total Environ* 2014; **472**: 862–71.
22. Nefau T., Karolak S., Castillo L., Boireau V., Levi Y. Presence of illicit drugs and metabolites in influents and effluents of 25 sewage water treatment plants and map of drug consumption in France. *Sci Total Environ* 2013; **461–462**: 712–22.
23. Banta-Green C. J., Field J. A., Chiaia A. C., Sudakin D. L., Power L., deMontigny L. The spatial epidemiology of cocaine, methamphetamine and 3,4-methylenedioxyamphetamine (MDMA) use: a demonstration using a population measure of community drug load derived from municipal sewage. *Addiction* 2009; **104**: 1874–80.
24. Thomas K. V., Bijlsma L., Castiglioni S., Covaci A., Emke E., Grabic R. *et al.* Comparing illicit drug use in 19 European cities through sewage analysis. *Sci Total Environ* 2012; **432**: 432–9.
25. Castiglioni S., Bijlsma L., Covaci A., Emke E., Hernandez E., Reid M. *et al.* Evaluation of uncertainties associated with the determination of community drug use through the measurement of sewage drug biomarkers. *Environ Sci Technol* 2013; **47**: 1452–60.
26. Lai F. Y., Ort C., Gartner C., Carter S., Prichard J., Kirkbride P. *et al.* Refining the estimation of illicit drug consumptions from wastewater analysis: co-analysis of prescription pharmaceuticals and uncertainty assessment. *Water Res* 2011; **45**: 4437–48.
27. United Nations, Department of Economic and Social Affairs, Population Division. World Urbanization Prospects: The 2011 Revision. CD-ROM edition. 2012. New York, NY.
28. European Monitoring Centre for Drugs and Drug Addiction. Purity of cocaine products at retail level, 2011. Available at: <http://www.emcdda.europa.eu/stats13#display:/stats13/ppptab7a> (accessed 12 December 2013).
29. Baker D. R., Barron L., Kasprzyk-Hordern B. Illicit and pharmaceutical drug consumption estimated via wastewater analysis. Part A: chemical analysis and drug use estimates. *Sci Total Environ*; in press; 2014; doi:10.1016/j.scitotenv.2013.11.107
30. Emke E., Evans S., Kasprzyk-Hordern B., de Voogt P. Enantiomer profiling of high loads of amphetamine and MDMA in communal sewage: a Dutch perspective. *Sci Total Environ*; in press; 2014; doi:10.1016/j.scitotenv.2013.11.043
31. European Monitoring Centre for Drugs and Drug Addiction. *EU Drug Market Reports—A Strategic Analysis*. Luxembourg: Office for Official Publications of the European Communities; 2013.
32. Griffiths P., Mounteney J., Lopez D., Zobel F., Götz W. Addiction research centres and the nurturing of creativity. Monitoring the European drug situation: the ongoing challenge for the European Monitoring Centre for Drugs and Drug Addiction (EMCDDA). *Addiction* 2012; **107**: 254–8.
33. Frischer M., Hickman M., Kraus L., Mariani F., Wiessing L. A comparison of different methods for estimating the prevalence of problematic drug misuse in Great Britain. *Addiction* 2001; **96**: 1465–76.
34. Ort C., Eppler J. M., Scheidegger A., Rieckermann J., Kinzig M., Sörgel F. Challenges of surveying wastewater drug loads of small populations and generalizable aspects on optimizing monitoring design. *Addiction* 2014; **109**: 472–81.
35. European Monitoring Centre for Drugs and Drug Addiction. *European Drug Report 2013: Trends and Developments*. Luxembourg: Office for Official Publications of the European Communities; 2013.
36. Pietsch J., Paulick T., Schulz K., Flössel U., Engel A., Schmitter S. *et al.* Escalation of methamphetamine-related crime and fatalities in the Dresden region, Germany, between 2005 and 2011. *Forensic Sci Int* 2013; **233**: 51–4.
37. Ort C., Lawrence M. G., Reungoat J., Mueller J. F. Sampling for PPCPs in wastewater systems: comparison of different sampling modes and optimization strategies. *Environ Sci Technol* 2010; **44**: 6289–96.
38. Ort C., Lawrence M. G., Rieckermann J., Joss A. Sampling for pharmaceuticals and personal care products (PPCPs) and illicit drugs in wastewater systems: are your conclusions valid? A critical review. *Environ Sci Technol* 2010; **44**: 6024–35.
39. Thomann M. Quality evaluation methods for wastewater treatment plant data. *Water Sci Technol* 2008; **57**: 1601–9.
40. O'Brien J. W., Thai P. K., Eaglesham G., Ort C., Scheidegger A., Carter S. *et al.* A model to estimate the population contributing to the wastewater using samples collected on census day. *Environ Sci Technol* 2014; **48**: 517–25.

Supporting information

Additional Supporting Information may be found in the online version of this article at the publisher's web-site:

Appendix S1 Questionnaire 2012 and 2013.

Appendix S2 Uncertainty estimation.

Appendix S3 Answers from questionnaire, all analytical data 2012 and 2013, means and *P*-values for changes of one week to another for all substances and locations (different spread sheets in separate Excel file).

Appendix S4 Weekly variation of drug loads 2012 (separate pdf file).