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Are cosmetics a significant source of PFAS in Europe? product inventories, chemical characterization and emission estimates†

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In this study, emission of per- and polyfluoroalkyl substances (PFAS) from the use of cosmetics in the European Economic Area (EEA; not including Lichtenstein and Iceland) was estimated for the first time. Using the European Commission database for information on cosmetic substances and ingredients (CosIng) ~170 structures containing at least $-CF_2-$ or $-CF_3$ were identified as ingredients in cosmetics on the European market. These structures were then cross referenced against the Cosmetic Database "CosMethics" to identify PFAS-containing products. Among these products, polytetrafluoroethylene (PTFE) and C9-15 fluoroalcohol phosphate were the most frequently listed PFAS ingredients. Thereafter, a sample of 45 cosmetics spanning 5 product categories was purchased in Sweden and characterized for total fluorine (TF), extractable organofluorine (EOF), and target PFAS. Using measured concentrations, the share of PFAS-containing products in each product category, sales data from Cosmetics Europe, as well as other parameters and assumptions, the annual emission of PFAS from cosmetics after use was estimated. Annual EEA-wide TF and EOF-based emissions ranged from ~17–38 000 kg F per year and 37–5100 kg F per year, respectively, representing combined emission to wastewater and solid waste (low to high emission scenario). Sum perfluoroalkyl carboxylic acid (PFCA) emissions were considerably lower (21 kg \sum PFCA per year; high scenario). While TF- and EOF-based emissions are significant, they are considerably lower than estimates of TF emission from washing of PFAS-coated textiles in the EU. This work provides the first estimate of PFAS emissions from cosmetics and highlights the importance of using a multi-platform analytical approach for PFAS emission estimates.

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Environmental significance

Per- and polyfluoroalkyl substances (PFAS) are listed ingredients in European cosmetic products, but their contribution to PFAS contamination in liquid and solid waste remains unclear. In this work the prevalence of PFAS in cosmetic products from the European market was investigated using a combination of database mining and comprehensive analytical characterization. These data, together with information on consumer habits related to cosmetic removal, were applied to estimate PFAS emissions to wastewater and solid waste for the first time. Since waste treatment does not remove PFAS completely, the disposal of PFAS-containing cosmetics may lead to environmental contamination. While PFAS emissions from cosmetics are significant, they are lower than emissions from other sources, such as washing of PFAS-coated textiles.

Introduction

Per- and polyfluoroalkyl substances (PFAS) are a diverse group of over 4700 chemicals, which have been widely used since the 1950s in industrial processes and consumer products.¹ The OECD/UNEP Global PFC Group defines PFAS as substances that

contain 1 or more fluorinated methyl or methylene carbon atoms (*i.e.* $-CF_3$ or $-C_nF_{2n-}$; $n \geq 1$), with some noted exceptions.² The extraordinary strength of the C–F bond together with the combined lipophobic and hydrophobic properties of perfluoroalkyl chains make PFAS useful for a wide variety of applications. Examples include non-stick cookware, textiles, food contact paper, medical equipment, motor oil fluids, fire-fighting foams, cosmetics and personal care products.^{1,3–5}

Since the first detection of PFAAs globally in humans and wildlife,⁶ concerns surrounding the risks of PFAS have steadily increased. PFAS have been associated with a wide range of health effects, including cancer, thyroid and hormonal disturbance, premature birth, neuro-developmental deficiencies and

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immunotoxicity.^{7,8} Phase-out and regulation of certain legacy PFAS such as perfluorooctane sulfonate (PFOS) and perfluorooctanoate (PFOA) has led to declining concentrations in humans from some parts of the world.^{9–11} However, some exemptions exist, and numerous PFAS remain unregulated and continue to be manufactured and used in consumer products globally.

Ingredients in cosmetics and personal care products are regulated by the European Parliament and Council of the European Union,¹² which stipulates that cosmetic manufacturers must ensure the safety of their products for human health. However, this regulation does not contain requirements on the use and release of substances that may impact the environment, such as PFAS. Instead, these substances are addressed by REACH, which stipulates that polymers and low molecular weight substances imported or manufactured in quantities of <1 tonne per year do not require hazard and risk assessments.¹³ PFOA and PFOA-related substances (*i.e.* substances that can degrade to PFOA) have been banned since the 4th of July 2020 in all products sold in the EU with limits of 25 and 1000 ng g⁻¹, respectively (REACH Annex XVII list of restricted substances (entry 68)). In addition, Sweden requires manufacturers and importers to register the occurrence of PFAS in products in the Swedish Chemicals Agency Products Register, regardless of the concentration used in the product. However, in other EU countries, such regulations are not in place and some PFAS may be completely overlooked in consumer products.

In addition to a variety of PFAS that are intentionally added to consumer products (mostly as emulsifiers, antistatics, stabilizers, surfactants, film formers, viscosity regulators and solvents), PFAS may occur unintentionally as impurities and/or degradation products.¹⁴ Fujii *et al.* (2013) were the first to measure PFAS in 24 cosmetics and personal care products, and found that 88% contained PFCAs, with sum PFCa concentrations up to 19 µg g⁻¹ in sunscreens and up to 5.9 µg g⁻¹ in other cosmetic products.¹⁵ Thereafter, Schultes *et al.* (2018) reported target PFAS in a wide range of cosmetic products from the Swedish market, showing high concentrations of PFCAs and PFCa-precursors in some samples, and even higher concentrations of total fluorine (TF) and extractable organofluorine (EOF).¹⁶ Most recently, a study of 231 cosmetics from Canada and the US reported that over 50% contained high levels of TF.¹⁷ While the possible occurrence of inorganic fluorine hampered determination of the exact proportion of PFAS-containing products, re-analysis of a subset of 29 products (*i.e.* ~13%) using targeted methods revealed detectable PFAS in all samples, with concentrations ranging from 22–10 500 ng g⁻¹ product weight.¹⁷ This is concerning, in particular since North American cosmetic producers are not necessarily required to list PFAS ingredients,¹⁷ thereby hampering consumer efforts to avoid PFAS-containing products.

Given the large surface area of the skin (~22 m²)¹⁸ and the liberal application of some cosmetic products, dermal uptake may represent an important route of exposure to PFAS in cosmetics.¹⁶ However, many PFAS in cosmetics are ionic or polymeric, both of which are expected to be poorly absorbed by

the skin.¹⁹ Dermal absorption of non-ionizable, low molecular weight PFAS is largely unknown. In addition to the potential for exposure from dermal uptake and ingestion of PFAS in cosmetics, PFAS used in these products may pose a risk to the environment. Environmental releases may occur during manufacture and use of cosmetics, or after use, when the products are intentionally removed from the skin or hair. These residues ultimately enter landfills or wastewater treatment plants (WWTPs) where they are often poorly retained and may subsequently enter the environment *via* landfill leachate,²⁰ WWTP effluent,²¹ or the use of sludge for agricultural fertilizer.²² However, to date, the prevalence of PFAS in cosmetics on large scale has not been determined; nor have emissions to waste streams.

The overarching aim of this work was to assess the importance of cosmetics from the European market as a source of PFAS to waste streams. To achieve this goal, an inventory of PFAS in cosmetics was carried out using European cosmetic databases. Thereafter, a selection of decorative cosmetics, hair care, and skin care products were characterized using a combination of targeted PFAS analysis and fluorine-specific measurements. Finally, using measured concentrations, the share of products containing PFAS, sales data, as well as other parameters and assumptions, the annual emission of PFAS into European wastewater and solid waste from cosmetics after use was estimated for the first time.

Materials and methods

Sample collection

An inventory of PFAS and their frequency of occurrence in cosmetic products was determined through database searches (Fig. 1). The European Commission database for information on cosmetic substances and ingredients (CosIng) was searched (for “fluoro”) to extract unique PFAS ingredients potentially occurring in cosmetic products using International Nomenclature of Cosmetic Ingredients (INCI) identifiers. All substances containing at least a –CF₂– or –CF₃ (including polymers), were included in the search. Note that substances considered “exceptions” (*i.e.* a carbon atom with a H/Cl/Br/I atom attached to it²) were also included. It is pertinent to note that one INCI name may comprise several PFAS (*e.g.* “C9–C15 fluoroalcohol phosphate”, seven PFAS) but for practical reasons, each INCI name was considered as a single PFAS for the database analysis.

The ~170 unique PFAS ingredients identified in CosIng were then cross referenced against the CosmEthics European database, in order to connect PFAS INCI names to specific cosmetic products within 11 product categories (Table S1†). Each of the product categories contained a range of sub-categories. For example, anti-aging cream, masks, and eye gel were among the sub-categories within the category “Facial Care” (Table S1†). In order to maximize sampling of sub-categories containing a high share of PFAS-containing products, the following approach was used: First, summary statistics on PFAS-containing products received from CosmEthics (EU/EEA barcode product database extract only) were reviewed and the share (%) of PFAS-containing products was calculated for each product sub-





Fig. 1 Schematic of cosmetic products database screening and selecting samples, analysis and emission estimation workflow. "PFAS-containing" and "PFAS-free" refer to the ingredient list on the product.

category. The most relevant sub-categories were identified as those with both the largest share of PFAS-containing products and the largest total number of products. Thereafter, a total of 50 anticipated samples was distributed among the most relevant product categories and sub-categories according to the number of PFAS-containing products in a given sub-category (*i.e.* percent distribution). Sampling aimed to collect the number of products identified from each of the targeted sub-categories (Table S1[†]).

In September 2020, a total of 45 different cosmetics were purchased (43 listing PFAS as ingredients, 2 which did not) either online (1 product) or from local stores in Stockholm. For further consideration and emissions calculations, products were re-categorized into the following 5 product categories: "decorative cosmetics", "hair care", "skin care", "toiletries", and "perfumes and fragrances". This was necessary in order to align with the categories of the retail sales data provided in a report by Cosmetics Europe²³ which was later used for emissions calculations (see below). Among purchased products, a total of 24 were "decorative cosmetics", 6 were "hair care" and 15 were "skin care" (out of which 2 products were for males). An overview of the investigated samples, brand names and listed fluorinated ingredients (*i.e.* INCI names) is available in Table S2.[†]

Chemical analysis

Standards and reagents. A total of 21 PFAS were targeted in this work. Linear isomer standards of perfluorobutanoate (PFBA), perfluoropentanoate (PFPeA), perfluorohexanoate (PFHxA), perfluoroheptanoate (PFHpA), PFOA, perfluorononanoate (PFNA), perfluorodecanoate (PFDA), per-

fluoroundecanoate (PFUnDA), perfluorododecanoate (PFDoDA), perfluorotridecanoate (PFTriDA), perfluorotetradecanoate (PFTeDA), perfluorohexadecanoate (PFHxDA), perfluorooctadecanoate (PFOcDA), perfluorobutane sulfonate (PFBS), perfluorohexane sulfonate (PFHxS), PFOS, perfluorodecane sulfonate (PFDS), perfluorooctane sulfonamide (FOSA), bis(1*H*,1*H*,2*H*,2*H*-perfluorooctyl)phosphate (6 : 2/6 : 2 diPAP), bis(1*H*,1*H*,2*H*,2*H*-perfluorodecyl) phosphate (8 : 2/8 : 2 diPAP) and (1*H*,1*H*, 2*H*,2*H*-perfluorooctyl) (1*H*,1*H*,2*H*,2*H*-perfluorodecyl) phosphate (6 : 2/8 : 2 diPAP) were purchased from Wellington Laboratories (Guelph, Canada). A list of isotopically-labelled internal standards is provided in Table S3.[†] Additional reagents used for sample preparation are provided in the ESI.[†]

Sample handling and instrumental analysis. All samples were analyzed directly (*i.e.* no sample extraction) by combustion ion chromatography (CIC) for determination of total fluorine (TF). In addition, a subset of 15 samples were selected in a manner which maximized the diversity of listed PFAS ingredients while covering the three main product categories and several subcategories. This subset was extracted with methanol and analyzed by CIC and liquid chromatography-tandem mass spectrometry (LC-MS/MS) for EOF and target PFAS, respectively. All extraction and instrumental analysis methods have been previously validated and reported (see Schultes *et al.* (2018)¹⁶ and also tables S3, S4, and text in the ESI[†] for details). The results of ongoing QA/QC procedures as part of this work are provided in the following section.



For comparison to TF and EOF data, targeted PFAS concentrations were converted to fluorine equivalents (C_{F_PFAS} ; ng F g^{-1}) using eqn (1):

$$C_{F_PFAS} = n_F \times A_F / MW_{PFAS} \times C_{PFAS} \quad (1)$$

where C_{PFAS} and n_F are the concentration and number of fluorine atoms for a given target, A_F is the atomic weight of fluorine (19.0 g mol^{-1}), and MW_{PFAS} is the molecular weight of the target (g mol^{-1}). Once the concentrations were converted to fluorine equivalents (*i.e.* ng F g^{-1}), they were summed to obtain ΣC_{F_PFAS} concentrations, which were directly comparable to EOF and TF measurements.

Quality assurance and quality control. Interlaboratory comparison of TF measurements performed at Stockholm University have shown good agreement with other labs for paper matrices.²⁴ In the present work, TF accuracy and precision were further confirmed through triplicate combustions of a certified reference material (CRM; BCR@-461, fluorine in clay), which was measured at the same time as real samples. The average percent recovery of these measurements was 90% (2.5% relative standard deviation [RSD], $n = 3$) and data were not corrected for losses. Boat blanks and a mid-level calibration standard were run intermittently to monitor potential carry-over and instrumental stability, respectively. All TF data were blank-subtracted and limits of detection (LODs) were calculated as 3 times the standard deviation of the blanks. Intra-sample variability of TF was assessed by analyzing 10% of samples in triplicate (*i.e.* 5 individual samples analyzed in triplicate).

Interlaboratory comparison of EOF measurements in water and sludge performed at Stockholm University have shown reasonable agreement with other labs.²⁵ For cosmetics, accuracy and precision were assessed by replicate matrix spike/recovery experiments, consisting of a PFAS-free cosmetic product which was extracted and analysed with and without fortification of a suite of PFAS (277 ng F fortification level; $n = 3$). The average percent recovery from these experiments was 61% (14% RSD) and data were not corrected for losses. Furthermore, an additional spiking experiment was performed using 250 ng sodium fluoride (NaF; $n = 3$), which was not recovered, confirming that inorganic fluorine was removed during the extraction procedure. In addition to analysing 10% of samples in triplicate (*i.e.* 2 individual samples analyzed in triplicate for EOF), three procedural blanks were handled in the same way as the samples to check for potential contamination introduced during the extraction procedure. A standard of PFOS (1 $\mu g mL^{-1}$) analyzed routinely with samples revealed good comparability to theoretical concentrations (92–113%), confirming that the response measured on the CIC was F-specific. Finally, potential carry-over and instrumental stability were monitored using the same procedure as explained for TF analysis. All EOF concentrations were blank-corrected and the limits of detection were calculated based three times the standard deviation of the procedural blanks.

For target PFAS measurements, accuracy and precision were assessed using the same replicate spike/recovery experiments as used for EOF, *i.e.* a PFAS-free cosmetic product which was

extracted and analyzed with and without fortification of 10 ng of individual PFAS (Table S3†). In addition to analyzing 10% of samples in triplicate (*i.e.* 2 individual samples analyzed in triplicate), three procedural blanks (*i.e.* no matrix) were handled in the same way as the samples to monitor for potential contamination introduced during the extraction procedure. Finally, during instrumental analysis, instrumental blanks and a mid-level calibration standard were run intermittently to monitor potential carry-over and instrumental stability, respectively. Procedural blanks did not show detectable contamination for any target PFAS. Therefore, limits of detection (LODs) were determined using the concentration obtained from the lowest calibration point with a well-shaped peak displaying an intensity >1000 and a signal-to-noise (S/N) ratio of >3 and converted to a weight-based concentration (*e.g.* ng g^{-1}) using the average sample weight (~0.1 g). Replicate spike/recovery experiments with target PFAS revealed an average percent recovery for all targets of 83% (RSD 13%; $n = 3$). Most substances fell within the range of 85–95%, with the exception of PFHxDA and PFOcDA, which had lower recoveries (61 and 36%, respectively). Considering that internal standards were added after extraction (necessary because a portion of the same extract was used for EOF analysis), these recoveries are reasonable, and reflect some losses incurred during the extraction procedure and/or matrix effects arising from the absence of exactly matching, isotopically labelled internal standards. Consequently, no correction for losses was made. Detection limits ranged from 1 to 18 ng PFAS g^{-1} cosmetic product sample, which are higher than for environmental sample analysis due to the low quantity of sample used.

As a final QC measure, a sample previously analyzed in Schultes *et al.* (2018) was re-analysed in the present study for TF, EOF and target PFAS concentrations. TF and EOF concentrations measured here were in good agreement with previous measurements (3100 *vs.* 2900 ng F g^{-1} and 1260 *vs.* 1380 ng F g^{-1} , respectively). For target analysis, sum PFCA (Σ PFCA) concentrations were also in good agreement (6.27 ng F g^{-1} *vs.* 8.11 ng F g^{-1} , respectively), but higher diPAP concentrations were observed in the present work compared to Schultes *et al.* (2018) (487 ng F g^{-1} here *vs.* 72 ng F g^{-1} in Schultes *et al.* respectively). This is perhaps not surprising considering that a different method was used for diPAP determination in Schultes *et al.* (2018) and because challenges were noted in the previous study for determination of PAPs in these samples due to their extremely high concentrations.

Emission estimates

Annual PFAS or fluorine emission (E_{PFAS} ; kg PFAS per year or kg F per year) was estimated after cosmetic use to wastewater and/or solid waste for the European Economic Area (EEA; not including Lichtenstein and Iceland, population roughly 450 million people) according to eqn (2).

$$E_{PFAS} = C_{PFAS} \times A_{products} \times f_{PFAS\ products} \times f_{release} \quad (2)$$

Each variable contains a number of calculations and assumptions which are explained in more detail in ESI (see



Section S3, Tables S8–S11†). C_{PFAS} is the concentration of PFAS (or fluorine) in a product obtained based on measurement of target PFAS (μg PFCAs per g product) or TF (μg F g^{-1} product), EOF (μg F g^{-1} product). The total amount of products sold annually (A_{products} ; tons per year) was derived from the retail sales price from the European cosmetic products market (EEA, not including Lichtenstein and Iceland), the market share of the different product categories,²³ and the assumed average product price and size per product category. The share of products containing PFAS ($f_{\text{PFAS products}}$; unitless) was derived by dividing the total number of cosmetic products containing PFAS by the total number of products for each product category (numbers from the CosmEthics database, but product subcategories recategorized into cosmetics Europe categories to match the sales statics categories, see Section S3, Table S12†). Finally, the fraction of the chemical released (f_{release} ; %) from the product into a certain compartment (*i.e.* wastewater or solid waste; see Section S3†) was obtained by considering consumer habits²⁶ and assumptions described in detail in the ESI (Section S3, Table S14†).

Finally, three different waste stream emission scenarios (*i.e.* low, average and high) were calculated for each of the three measurements (*i.e.* TF, EOF and targeted PFAS) based on: (i) concentrations (minimum, average and maximum, Tables S5–S7 and explanation in ESI†) in each product category; (ii) three release scenarios (*i.e.* f_{release} % into wastewater and solid waste, Table S14†) based on statistics on consumer removal habits²⁶ (Table S13†) and (iii) three product price levels (Table S10†), for the overall product categories. The difference between the total amount of PFAS in cosmetics and the amount that is emitted *via* wastewater was considered to end up in solid waste (*i.e.* disposal of cosmetic products removal pads/tissues *etc.* into waste, left-over product amount in the package, disposal after a product's lifetime). Further, PFAS emissions during the production of cosmetic products are not considered, neither are releases of volatile PFAS into the air. A detailed explanation of each input parameter for the different emission scenarios is provided in the ESI,† as well as an overview table on the ingoing parameters and their values applied in the emission calculation for the different scenarios (Tables S15–18†).

Result and discussion

Product characterization

TF and EOF analysis. TF was measured in a total of 45 cosmetics (43 PFAS-containing products and 2 not listing PFAS ingredients; Fig. 2 and Table S5†). The 43 PFAS-containing samples comprised of 23 decorative cosmetics, 14 skin care and 5 hair care products. Among products listing fluorinated ingredients, TF concentrations ranged from below LOD (*i.e.* <0.001 mg F g^{-1} in one hair spray sample and <0.035 mg F g^{-1} one mask) to 13.8 mg F g^{-1} sample (one skin care product). TF concentrations in skin care products were the most variable ($<\text{LOD}$ – 13.8 mg F g^{-1}), followed by decorative cosmetics (0.02–6.01 mg F g^{-1}) and hair care ($<\text{LOD}$ – 0.05 mg F g^{-1}). Considerable variability was also observed within sub-categories, *e.g.* blusher/bronzer/contour (0.90–5.14 mg F g^{-1}), foundation/BB

cream (0.02–3.31 mg F g^{-1}), anti-age cream (0.08–3.80 mg F g^{-1}), and mask ($<\text{LOD}$ – 10.6 mg F g^{-1}). Generally, the TF range of the cosmetic products compared well to prior studies on cosmetic products in Sweden¹⁶ (<0.091 –19.2 mg F g^{-1}), Denmark²⁷ (0.03 to 0.740 mg F g^{-1}), and the US¹⁷ (0.155–3.96 mg F g^{-1}).

As was shown in Schultes *et al.* (2018), there was some evidence that inorganic fluorine contributed to the TF signal in some samples.¹⁶ For instance, TF concentrations were above the LOD for eye liner pen 1, which did not list any PFAS ingredients. Assuming that the fluorine does not occur from a residual or undeclared PFAS ingredient, it is likely attributable to inorganic substances (*e.g.* synthetic fluorophlogopite/mica or fluoride). Indeed 24 samples listed “mica” (not specified as synthetic or natural) and 7 samples listed “synthetic fluorophlogopite” (Fig. 2 and 3) but the exact quantities remain unclear. While EOF and targeted PFAS analysis do not capture inorganic fluorine and can be used to rule out its occurrence (as was the case here), these approaches may underestimate the total PFAS concentration (depending on the PFAS present), since some organofluorine substances may not be extracted with the solvent and/or measured in either case (*i.e.* during EOF or targeted PFAS analyses). We conclude that all three approaches may be susceptible to bias (TF – over-reporting; EOF and targeted PFAS – under-reporting); therefore, a combined approach, presenting upper and lower bound estimates, is the most transparent and comprehensive strategy for product characterization.

EOF measurements were performed in 15 of the 45 cosmetic products. Concentrations of EOF ranged from $<\text{LOD}$ (<162 ng F g^{-1} for exfoliator, <325 ng F g^{-1} for concealer 2 and Eyeshadow 5) up to 4.93 mg F g^{-1} for foundation/BB cream 4 (Fig. 3a, Table S6†). The second highest EOF concentration was detected in foundation/BB cream 3 (1.58 mg F g^{-1}). Similar to TF, EOF concentrations varied both within product categories and sub-categories. While skin care samples appeared to have lower EOF concentrations than hair care and decorative cosmetic products, the small sample size ($n = 6$) made it difficult to draw firm conclusions about trends of PFAS concentrations among product types. Overall these results compared well with prior work by Schultes *et al.* (2018) which reported EOF concentrations of $<\text{LOD}$ up to 1.72 mg F g^{-1} , with the exception of one product in the current study which contained EOF concentrations up to 4.93 mg F g^{-1} (foundation/BB cream 4).

For most samples, EOF concentrations were ~ 10 – $10\,000$ -fold lower than TF concentrations, indicating that the listed PFAS ingredient(s) were poorly extracted and/or that the products contained high concentrations of inorganic fluorine (Table S16†). The few instances of EOF exceeding TF concentrations (*e.g.* foundation/BB creams 3 and 4, pressed powder 2) are attributable to the variability in TF and EOF analyses, and have been documented previously.²⁸ In these three cases the EOF and TF are considered equivalent, indicating that the organofluorine ingredient is efficiently extracted and that inorganic fluorine is negligible (Table S16†). However, the observation of EOF concentrations at similar levels to TF in the case of pressed powder 2, was surprising, considering that the listed ingredient (polyperfluoroethoxymethoxy difluoroethyl PEG phosphate)





Fig. 2 TF concentrations ($\mu\text{g F g}^{-1}$) of different cosmetics, grouped according to product category. Bar colors represent samples containing polymers (red), perfluorinated cyclic/aliphatic species lacking a functional group (yellow) and PFAS species with functional group(s) (grey) that appeared on the ingredient list. Samples of mask 1, treatment 2 and hairspray 2 are < LOD. Structures T and Y are adopted from literature,²⁹ but are uncertain based on the INCI name provided. Samples labelled PFAS-FREE did not list any PFAS ingredients. *Mica listed among ingredients; #synthetic fluorophlogopite listed among ingredients.

appears to be polymeric and was not expected to be extracted efficiently with methanol. In comparison, all other polymer-containing products (e.g. Eyeshadow 5 and lip liner, pen 1, eye liner, pen 2 and aftershave) displayed EOF levels that were much lower than TF (range 0.01–6.8%). It can be speculated that the polymer in pressed powder 2 is much smaller (i.e. shorter chain length) and more soluble in methanol than the two other polymers in the aforementioned products.

Targeted analysis. Target PFAS measurements were performed for the same samples subjected to EOF analysis (i.e. 15 of the 45 cosmetic products); and the results of these analyses are summarized in Fig. 3b. In total, 7 out of 15 samples contained measurable concentrations of at least one PFAS. \sum PFCA concentrations ranged from <LOD to 9560 ng g^{-1} with the highest concentrations in mask 2. The second highest \sum PFCA concentration (425 ng g^{-1}) was observed in another mask product (mask 1, Table S7†). Samples of foundation/BB creams, masks, concealer 1, eye shadow 5, lip liner pen 1 contained up to 7 out of the 21 monitored PFAS above detection limits, whereas samples of anti-age cream 2, exfoliator, lip liner pen 2, pressed powder 2, concealer 2, shampoo, styling cream and aftershave did not contain detectable concentrations of targeted PFAS. None of the PFASs from the target list were detected above the LOD, which is in accordance with previous studies from

Europe,^{14,16} but contrasts recent work on North American cosmetic products, which detected low (i.e. < 30 ng g^{-1}) concentrations of PFASs in most Canadian products and 1 US product. The most frequently detected PFCA was PFBA in seven of the products, followed by PFHxA in three of the products and PFHpA in two of the products. PFPeA, PFOA, PFNA, PFDA, PFDoDA and PFTeDA were each detected in one sample (PFPeA in foundation/BB cream 4, the other PFCAs in foundation/BB cream 3, Fig. 3b). PFOA was measured in one sample and occurred at concentrations above the EU limit of 25 ng g^{-1} (112 ng g^{-1} in foundation/BB cream 3).³⁰ Notably, analysis of 29 North American cosmetic products¹⁷ revealed sum PFAS concentrations ranging between 22 and 10 500 ng g^{-1} , with 6 : 2 fluorotelomer alcohols (FTOHs) and methacrylates (FTMAs) among the dominant PFAS. While FTOHs and FTMAs were not included in the present study, it was expected that both classes of PFAS (if present) would be captured by EOF measurements. Overall, the diversity of PFAS ingredients and residuals in consumer products which are not captured by targeted approaches highlights the importance of including TF- and EOF-based measurements for a comprehensive picture of PFAS contamination in consumer products.

Our targeted PFAS analysis also detected intentionally added PFAS ingredients in two products. Foundation/BB cream 3 and 4





Fig. 3 (a) Fluorine mass balance showing TF, EOF and targeted PFAS concentrations ($\mu\text{g F g}^{-1}$ sample). The targeted PFAS ($\sum\text{PFAS}$) concentrations were converted to fluorine concentrations for comparative purposes. ($\text{LOD}_{\text{TF}} = 35.45 \mu\text{g F g}^{-1}$, $\text{LOD}_{\text{EOF}} = 0.32 \mu\text{g F g}^{-1}$, $\text{LOD}_{\sum\text{PFAS}} = 0.0015 \mu\text{g F g}^{-1}$). *Mica listed among ingredients; #synthetic fluorophlogopite listed among ingredients. (b) Concentrations (ng g^{-1} sample) of targeted PFAS in 5 decorative cosmetics and 2 skin care samples (masks). Samples not shown (8 out of 15) were <LOD. Samples are plotted on a logarithmic scale (base 10) to enhance readability. The letters on top of each bar correspond to structures provided in Fig. 2.

contained “perfluoroalkylethyl phosphate” and “fluoroalcohol phosphate”, respectively, which assumably refers to polyfluoroalkyl phosphate esters (PAPs). Indeed, diPAPs were measured in both products, at concentrations up to 2.3 mg g^{-1} (6 : 2 diPAP; Fig. 3b). 6 : 2/8 : 2 and 8 : 2 diPAP were detected at concentrations up to 0.67 mg g^{-1} (Fig. 3b), which far exceeds the EU limit of 1000 ng g^{-1} for PFOA-related substances.³⁰ These diPAP concentrations exceeded concentrations of all other PFAS by several orders of magnitude in some cases, demonstrating that targeting listed ingredients will greatly increase $\sum\text{PFAS}$ concentrations, compared to if exclusively impurities are measured (*e.g.* PFCAs, FTOHs, *etc.*). This underlines the need for more analytical standards and methods in order to quantify the diversity of PFAS which are intentionally added to products.

PFAS emissions

Low, high, and average emission scenarios for PFAS from the use of cosmetics were estimated using TF, EOF, and $\sum\text{PFCA}$ ($\text{C}_4\text{--C}_{18}$)-concentrations, resulting in a total of 9 estimates for combined emission to wastewater and solid waste (Table 1).

These estimates should not be confused with environmental emissions, since the fraction of PFAS retained and/or destroyed during treatment (*i.e.* by wastewater treatment plants, landfills, or incineration plants) has not been included in the calculation, nor were emissions occurring during cosmetic production. For TF-based estimates, emissions ranged from 17 kg F (low emission scenario) to $38\,000 \text{ kg F}$ (high emission scenario) per year. Since TF concentrations capture all fluorinated substances (both organic and inorganic) these estimates will err on the side of overestimation. Moreover, caution is warranted when converting these fluorine concentrations to PFAA-equivalents (for example to estimate surface water concentrations), since many of the fluorinated ingredients captured by TF measurements are not PFAA-precursors and may not be water soluble. Rather, they are polymers which may contribute to particulate (*i.e.* nano- or microplastic) pollution, or possibly retained in the wastewater treatment plant. By comparison, EOF-based emission estimates, which ranged from 37 kg F (low emission scenario) to 5100 kg F (high emission scenario) per year do not include contributions from inorganic fluorine, but are limited to (semi-) polar PFAS, typically PFAAs and/or PFAA-precursors. Polymeric



Table 1 Annual estimates for total (emissions to wastewater and solid waste), each in a low, average and high emission scenario for the different cosmetic product categories based on the TF (kg F per year), EOF (kg F per year) and \sum PFCAs (kg \sum PFCAs/per year) measurements. All values are rounded to two significant figures. Emission estimates for Perfumes and Fragrances category were negligible and are not shown. The higher emissions for EOF compared to TF (low emission scenario) is because the lowest measured EOF concentration among 15 samples characterized for EOF was higher than the lowest measured TF concentration among the 45 samples characterized for TF. A value of "0" indicates negligible emission for a given emission scenario based on the input parameters and assumptions

Product category	Emission								
	Low emission scenario			Average emission scenario			High emission scenario		
	Total (Wastewater + solid waste)	Wastewater	Solid waste	Total (Wastewater + solid waste)	Wastewater	Solid waste	Total (Wastewater + solid waste)	Wastewater	Solid waste
TF (kg F per year)									
Skin care	8.0	5.3	2.7	8200	6200	2000	29 000	29 000	0
Toiletries	1.0	0.82	0.21	560	500	58	1500	1500	0
Hair care	1.9	1.5	0.35	1000	930	86	2700	2700	0
Decorative cosmetics	5.6	1.0	4.6	1200	650	570	4100	4100	0
Cosmetics total	17	8.6	7.9	11 000	8300	2700	38 000	38 000	0
EOF (kg F per year)									
Skin care	0.17	0.11	0.059	11	8.7	2.8	78	78	0
Toiletries	13	10	2.7	310	270	32	570	570	0
Hair care	24	20	4.4	560	510	47	1000	1000	0
Decorative cosmetics	0.11	0.020	0.092	380	200	180	3400	3400	0
Cosmetics total	37	30	7.3	1300	1000	260	5100	5100	0
\sumPFCAs (kg \sumC₄-C₁₈ PFCAs per year)									
Skin care	0	0	0	2.7	2.02	0.64	20	20	0
Toiletries	0	0	0	0	0	0	0	0	0
Hair care	0	0	0	0	0	0	0	0	0
Decorative cosmetics	0	0	0	0.035	0.019	0.016	0.24	0.24	0
Cosmetics total	0	0	0	2.7	2.0	0.66	21	21	0

PFAS and low molecular weight non-polar PFAS are unlikely to be captured in these estimates. Finally, \sum PFCAs-based estimates, which are expressed on a kg \sum C₄-C₁₈ PFCAs per year basis (instead of fluorine equivalents per year) ranged from negligible (low emission scenario) to 21 kg \sum C₄-C₁₈ PFCAs per year (high emission scenario), and were dominated by emissions to wastewater in all 3 scenarios. These estimates represent an even more severe underestimate of total PFAS emissions, since only PFCAs are included, yet they provide a good estimate of contamination introduced from unintentionally added residual PFAAs and can be more readily used to estimate contributions to PFAS concentrations in surface water.

To investigate the importance of PFAS emission from cosmetics relative to other sources, emission estimates calculated here were compared to literature data, expressed on a fluorine equivalent basis (Fig. 4). While cosmetics after use release up to 0.015 tonnes F from C₄-C₁₈ PFCAs annually within the EEA, annual emission of C₄-C₁₄ PFCAs during the entire life-cycle of PFOA and PFNA-based products between 1951 and 2015 were much higher (25–240 tonnes F per year).³¹ In comparison, Li *et al.* (2017) estimated that degradation of side-chain fluorotelomer-based polymers in waste stocks would release ~34–635 tonnes of C₄-C₁₂ PFCAs per year from 2015

onward into the environment³² (Fig. 4). Considering PFCAs emissions alone, the contribution from cosmetics appears low. However, it is pertinent to note that our emission estimates for cosmetics only include release from the products themselves, whereas other studies included the emission from production as well. Considering the high emission scenario using TF data, emissions from cosmetics are only 3.5-fold lower than emissions from outdoor textile washing (*i.e.* 38 tonnes F per year *vs.* 133 tonnes F per year)³³ and TF data are in the same range as estimates for the global emissions of fluorine during the life cycle of PFHxS- and PFDS-based products (0.4–11.1 tonnes F per year compared to the cosmetic range 0.02–38 tonnes F per year).³⁴

Implications

The present work demonstrates that both regulated and unregulated PFAS exist in cosmetic products from the EEA, sometimes at concentrations exceeding EU limits for PFOA (25 ng g⁻¹) and PFOA-related substances (1000 ng g⁻¹) since the 4th of July 2020 for products sold in the EU.³⁰ Considering that regulation of PFOA and PFOA-related substances under REACH has existed since 2019 and are now included under Annex A of the Stockholm Convention, new products or product version





Fig. 4 Comparison of PFAS emission from other consumer products. Values in the brackets show the fluorine equivalent emission calculated for each study; ranges represent low to high emission scenarios. Note that cosmetics emissions only consider PFAS release after the use of cosmetics, while emissions from the life cycle of PFDS- and PFHxS-based products considers production as well.

ought to exclude these compounds. While the proportion of products listing PFAS ingredients is small relative to the total number of products on the market, the emissions to wastewater and solid waste may be significant, albeit lower than other sources (*e.g.* outdoor textiles). Nevertheless, questions remain as to why these products are needed in cosmetics or why some regulated PFAS still appear in these products and how to exclude them in case of impurities.

Study limitations, uncertainties and future improvements

This study was subject to a number of limitations and uncertainties related to inventory development, product sales estimates, analytical characterization and emission estimates, which are described in detail in the ESL.[†] A brief overview of the most important sources of error from each parameter is described here:

Inventory development. The total number of PFAS occurring in cosmetic products and/or existing as INCI names is likely an underestimate, either because products were missed during the database search, were not named correctly in the database, or because one INCI name often includes several different PFAS.

Analytical characterization. A weakness of the study was that the total number of samples analyzed represents a small fraction of the total number and variety of products available. Moreover, the focus of the study was primarily on products with listed PFAS ingredients; the extent of PFAS contamination in supposedly “PFAS-free” (according to listed ingredients) products in the EEA remains unclear. While high PFAS concentrations were previously observed in North American cosmetics¹⁷ which did not disclose PFAS ingredients, a similar result in European products would be surprising, given the requirements

for disclosure of ingredients. However, PFAS as impurities from the raw materials or subsidiary technical materials cannot be ruled out in the current study either, as they do not count as ingredients (Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products). Neither can PFAS be ruled out in products that do not follow the existing legislations, which we discovered in a few cases.¹² Further, emissions based on TF, EOF, or target PFAS each have advantages and limitations: while TF captures all PFAS it may be subject to overestimation due to the co-occurrence of inorganic fluorine. On the other hand, while EOF is not affected by inorganic fluorine (assuming an appropriate extraction procedure is used to remove it), some PFAS are clearly not extractable using the polar solvents employed here. Thus, EOF may underestimate the PFAS content in some cases. Finally, target PFAS analysis is useful for sensitive and quantitative determination of individual substances, but undoubtedly underestimates total PFAS content, since listed PFAS ingredients are seldom included in targeted analytical methods and since the PFAS analyzed for was in this study limited to PFCAs, PFASs and PAPs.

Emission estimates. PFAS emissions estimated in the present work were only calculated for the product after use and release to wastewater and solid waste. The releases during production and potential losses while use (*e.g.* to air) was not considered, nor the fraction released to the environment following treatment of solid or liquid waste. Finally, information on consumer habits pertaining to cosmetic removal were not available for all product categories, thus assumptions were required. Since the quantity of cosmetic products sold per year (on a mass basis) was unavailable, several assumptions were



made in order to estimate these values, which could lead to large differences in emission estimates. The two parameters with the biggest influence on the cosmetic product amount sold per year are likely the price per product and the size of a product. A 10% change in either of these parameters would each result in a corresponding 10% change in the total emission estimates.

Conflicts of interest

There are no conflicts to declare.

Preprint

A preprint version of this work was published as a government report (<https://www.kemi.se/en/publications/pms/2021/pm-9-21-pfass-in-cosmetics>).

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