Per- and polyfluoroalkyl substances (PFASs) in Swedish household dust and exposure of pet cats

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Abstract
Per- and polyfluoroalkyl substances (PFASs) are used in a wide range of products and have been found ubiquitously in our indoor environment, and there is evidence that exposure to PFAS can lead to adverse endocrine effects, such as thyroid hormone disruption. Pet cats have a high dust intake due to their grooming behavior and have been shown to be a suitable sentinel species for assessment of toddler’s exposure. Here we used paired household dust (n=46) and cat serum (n=27) samples to establish whether dust is a relevant exposure pathway to PFASs. An analytical method for PFAS analysis was optimized using a low volume of cat serum samples, combining solid-phase extraction and online sample cleanup. Dust was extracted with methanol by sonication and cleaned up by addition of active carbon. In total, 27 PFASs were analyzed by liquid chromatography/mass spectrometry analysis. The correlation between PFAS levels in dust and serum, serum lipids and thyroid hormone levels, and PFAS levels in dust between different rooms were statistically evaluated. PFOS and PFDA could be quantified in all cat serum samples (median 2300 pg/mL and 430 pg/mL, respectively), followed by PFOA (median 1100 pg/mL), quantified in 96% of the samples. The levels of 6:2 and 8:2 diPAPs were determined in 65% and 92% of the serum samples, respectively, and were an order of magnitude lower (1.4–160 pg/mL). Household dust on the other hand was dominated by 6:2 and 8:2 diPAPs, with a median of 65 ng/g dust and 49 ng/g dust, respectively. PFOS (median 13 ng/g dust) and PFOA (median 9 ng/g dust) were quantified in 93% of the dust samples. Only eight PFASs were detected (LOD) in at least 50% of the samples of both matrices and could be paired. Significant correlations between cat serum and dust were found for PFOA (rs=0.32, p<0.049) and PFUnDA (rs=0.55, p<0.001). Significant positive correlations were found between serum total thyroxine (rs=0.11, p<0.05) and PFNA and between serum cholesterol and PFHpA (rs=0.46, p<0.01), PFUnDA (rs=0.40, p<0.05), PFDaDA (rs=0.44, p<0.01), and sum PFAS (rs=0.48, p<0.01). In conclusion, this study confirmed that dust is a relevant exposure pathway for the ingestion of some PFASs for cats, and the serum levels of PFASs could be of relevance for the cat’s health.

Keywords PFAS · Dust · Cats · Serum · Thyroid · PFOS · PFOA · PAPs

Introduction
It has been shown over the last decades that anthropogenic organohalogen contaminants (OHCs) may interact with hormones and act as endocrine disrupting compounds (EDCs) in wildlife and humans, leading to various diseases, such as diabetes and cancers related to the endocrine system (WHO/UNEP 2013). In addition, exposure to EDCs may lead to the onset of immune-related, thyroid-related, and neurodevelopmental disorders (WHO/UNEP 2013). Thyroid hormone disorders have gained increased attention as evidence supporting the disruption of thyroid hormone mechanisms in humans and wildlife during the sensitive embryonic and postnatal development have been repeatedly reported (WHO/UNEP 2013). There are several
Pathways through which organisms may be exposed to EDCs, with food being one of the major routes for OHCs. Another significant exposure route is dust ingestion, especially for toddlers and children at developing stages due to their extensive hand-to-mouth activity and low body weight. For modeling purposes, the estimated daily intakes of dust ingested are between 40 and 100 mg for children and 20–60 mg for adults (US-EPA 2017; Jones-Otazo et al. 2005).

Domestic pet cats have been used as a sentinel species for exposure to OHCs accumulated in dust in indoor home environments due to their high dust intake related to their natural grooming behavior (Ali et al. 2013; Bost et al. 2016; Chow et al. 2015; Dírku et al. 2013; Dye et al. 2007; Guo et al. 2016; Henríquez-Hernández et al. 2017; Mizukawa et al. 2016; Mizukawa et al. 2017; Norrgran Engdahl et al. 2017; Walter et al. 2017; Wang et al. 2018). To our knowledge, there are no studies estimating the daily intake of dust for cats, although correlation between levels of OHCs in dust and cat serum has been confirmed (Norrgran Engdahl et al. 2017). In addition, cats have an increasing prevalence of developing hyperthyroidism, an endocrine disorder also increasing in humans, and associations have been found between elevated levels of certain polychlorinated dibenyl ethers (PCBs) in hyperthyroid cats compared to levels in healthy cats (Norrgran et al. 2015). Feline hyperthyroidism is manifested as a disturbed upregulation of the thyroid hormone (TH), thyroxine (T4) concentration, and downregulation of thyroid stimulating hormone (TSH), with clinical consequences such as weight loss, polyphagia, polydipsia, hyperactivity, aggression, diarrhea, vomiting, and tachycardia (Jones et al. 2019; Peterson and Ward 2007).

In most organisms, THs are transported, bound to the blood serum proteins albumin (ALB), transthyretin (TTR), and thyroxine-binding globulin (TBG) (Richardson 2009). Only TTR and ALB act as TH-binding transport proteins in cats, and TBG has not been found in cat serum (Larsson et al. 1985). ALB is a general transport protein, which occurs in an excessive amount in blood plasma, but ALB has a low specific binding of THs. The transport protein TTR, on the other hand, is highly specific and can transport THs across membranes, such as the blood-brain barrier, across the placenta and into cerebrospinal fluid (Meerts et al. 2002; Schreiber et al. 1995; White and Kelly 2001). Several anthropogenic OHCs have been demonstrated to be able to bind to TTR and competitively inhibit the transport of T4 (Weiss et al. 2015). The majority of the compounds reported to be TTR binders are aromatic and hydroxylated and contain halogens, features they have in common with T4 itself. One compound group that differs structurally but has been shown to bind to TTR are per- and polyfluoroalkyl substances (PFASs) (Weiss et al. 2009).

PFASs exhibit high surface activity due to their hydrophobic and hydrophilic characteristics, and the many carbon-fluor bonds make them extremely stable. This gives them useful properties, e.g., thermal stability, acid resistance, and ability to lower surface tension. Due to this, PFASs can be found in a wide range of industrial and commercial products, such as polymers, surfactants, lubricants in textile coatings, non-stick coatings, stain repellents, food packaging, and firefighting foams (Buck et al. 2011; Wang et al. 2017). The human exposure pathways are mainly via food and drinking water, although PFASs can also diffuse directly from applied household products and associate to dust particles (Vestergren and Cousins 2009). Recent studies have indicated a correlation between concentrations of PFASs in household dust and human serum, particularly for small children (Beesoon et al. 2012; Wu et al. 2015). PFASs have been determined in cat serum from the USA at levels similar to the US population based on the NHANES (National Health and Nutrition Examination Survey) database (Bost et al. 2016; Wang et al. 2018).

The aim of this study was to determine the PFASs used in common consumer products in paired samples of household dust and serum of domestic cats from a central region of Sweden. The samples analyzed are part of a larger project that aimed to identify and evaluate mixtures of thyroid hormone disruptors in our indoor environment (MiSSE—Mixture Assessment of EDCs, Formas 210-2012-131). The PFASs analyzed are perfluoroalkyl carboxylic acids (PFCA), perfluoroalkyl sulfonic acids (PFSA), perfluoroalkyl phosphonic acids (PFPA), perfluoroalkyl phosphinic acids (PFPIA), polyfluoroalkyl phosphate mono- and diesters (PAP and diPAP), 6:2 fluorotelomer sulfonic acid (6:2 FTSA), and perfluorooctane sulfonamide (FOSA).

**Materials and methods**

**Samples**

Seventeen families in Stockholm and Uppsala region (Sweden) participated in the study. Samples were obtained between August 2013 and March 2014. The inclusion requirements of families included having a healthy pet cat and at least one child under 10 years old living at home. The details of participating households and their cats have been reported elsewhere (Norrgran Engdahl et al. 2017). In total, blood was drawn from 29 unsedated pet cats in their home environment. Serum from two pairs of siblings from two homes was pooled due to low sample volume. Thus, 27 cat serum samples (25 individual samples and two pooled samples) from 17 homes are reported. On average, families had two children living at home (1–13 years old) and more than one cat. The majority of families lived in free-standing houses (53%). The majority of cats (86%) spent more than 50% of their time...
indoor. The cats’ health was checked by a simple clinical examination before blood was sampled. Their thyroid status was evaluated by measuring levels of total serum T4 and TSH at the laboratory of the university animal hospital at the Swedish University of Agricultural Sciences. All cats were clinically healthy and not on medication.

Blood was taken from the cephalic vein in the right foreleg with a 0.8-mm (21 G) needle into plain evacuated tubes with a cloting activator (BD Vacutainer, BD, Plymouth, UK). Serum was obtained by letting whole blood coagulate at ambient temperature for at least 30 min and then centrifuged for 5 min (3000 G), and the supernatant was collected and stored at −20°C prior to analysis.

Blood lipid content was determined enzymatically for triglycerides and cholesterol. Total lipid was calculated using an average molecular weight for triglycerides and cholesterol of 807 and 571 g/mol, respectively, and assuming free and esterified cholesterol in a ratio of 1:2 (Covaci et al. 2006). Triglycerides and cholesterol were measured at the laboratory of the university animal hospital at the Swedish University of Agricultural Sciences (Norrgran Engdahl et al. 2017).

Still-standing dust was collected using a dust collector (Dustream®, Indoor Biotechnologies Ltd., Wiltshire, UK) containing a disposable filter (mesh size 40 μm) attached to a household vacuum cleaner tube. Rooms sampled were the living room, the adult’s bedroom, the child’s room, and if available, an additional playroom of the participating families. The samples were sieved (1 mm) to isolate the estimated ingested fraction for cats (0.04–1.0 mm). Larger particles such as food crumbs, gravel from outside, or hair were avoided. Due to limited sample amount and other chemical analysis performed within the project, not all of the rooms from the 17 families could be analyzed for PFASs. Dust samples from the following rooms were available: child’s room, adult’s bedroom, living room, and additional play room (n=5). In total, 46 dust samples were analyzed, representing all 17 homes.

Chemical analysis

All PFAS abbreviations are taken from Buck et al. 2011 (Buck et al. 2011). The internal standards (IS) used for quantifying dust and serum levels, with the corresponding native compounds quantified with each isotope in brackets, are 13C4-PFBA (PFBA), 13C5-PFPeA (PFPeA), 13C2-PFHxA (PFHxA), 13C4-PFHpA (PFHpA), 13C4-PFOA (PFOA), 13C5-PFNA (PFNA), 13C2-PFDA (PFDA), 13C2-PFUnDA (PFUnDA), 13C2-PFDoDA (PFDoDA, PFTrDA, PFTeDA), 18O2-PFHSs (PBBS, PFHxs), 13C2-PFOS (total PFOS, i.e., branched and linear), 13C2-6:2 FTSA (6:2 FTSA), 13C4-6:2 diPAP (6:2 diPAP, 6:6 PFPIA, 6:8 PFPIA), 13C4-8:2 diPAP (8:2 diPAP, 8:8 PFPIA), 13C2-6:2 PAP (6:2 PAP), 13C2-8:2 PAP (8:2 PAP), and Cl-PFHxPA (PFHxA, PFPOA, PFDA). 13C3-PFHxA and 13C8-PFOA were used as recovery injection standards. All analytical standards were purchased from Wellington Laboratories. The calibration curve (concentration range 0.02–5 ng/mL), IS mixture, and injection standards were prepared in methanol (MeOH, J.T. Baker). The PFSA, PFCA, 6:2 FTSA and FOSA, and the organophosphorus PFASs (OP-PFASs; PAPs and diPAPs, PFPeAs, and PFPIAs) needed different sample pre-treatment and chemical analysis settings, specified below.

Serum

Fifty microliter cat serum was used for the PFSA, PFCA, 6:2 FTSA, and FOSA analysis. Before sample treatment, the IS mixture (50 μL, concentration 10 ng/mL) and MeOH (150 μL) were added. A larger volume of cat serum (120–520 μL) was used for the OP-PFAS analysis; the IS mixture (50 μL) and MeOH with 1% formic acid (400 μL, Sigma-Aldrich) were added. All samples were vortexed 30 s and sonicated 10 min, followed by centrifugation (13000 rpm) for 10 min. The supernatant was transferred to a vial, and the whole sample was injected into an online system containing a 5 μm C8 solid-phase extraction (SPE) column (4.6 * 10 mm, XTerra MS Waters). The details of the solvent and pump conditions are given in supporting information in Additional file 1 (Table SI-2).

Dust

The IS mixture (50 μL, concentration 20 ng/mL) was added to the sieved dust (50 mg). The sample was extracted twice with 2 mL followed by 1 mL MeOH, with 30-s vortex and 15-min sonication in between. The combined extract was centrifuged (3000 rpm) for 10 min, and the supernatant was transferred into a new tube. The extract (3 mL) was split into two equal portions for separate cleanup procedures for analyses of PFSA, PFCA, 6:2 FTSA and FOSA, and OP-PFASs. The extract for PFSA, PFCA, 6:2 FTSA, and FOSA analyses was diluted in 25 mL water (Milli-Q, obtained from a Milli-Q Reference A+ purification system, Millipore, Bedford, MA, USA), and the pH level was adjusted with ammonium hydroxide (NH4OH, Sigma-Aldrich) to pH 10. For liquid/liquid extraction, hexane (4 mL, J.T. Baker) was added to the water, mechanically shaken for 10 min, and subsequently the two phases were separated by centrifugation (3000 rpm). The organic layer was discarded. The pH of the water phase was adjusted to pH 4 by adding hydrochloric acid (Sigma-Aldrich). For cleanup, SPE cartridges (Oasis WAX 150 mg 6cc, Waters) were used. After conditioning the cartridges and loading the extract, the cartridges were washed with 4 mL ammonium acetate (NH4Ac, pH 4, Sigma-Aldrich) and 8 mL tetrahydrofuran (Biosolve/MeOH (3/1, v/v)), and the extract was eluted with 4 mL methanol 0.1% NH4OH. The eluate was evaporated to dryness and reconstituted in 200 μL 50/50 MeOH/Milli-Q water (v/v).
The extract for OP-PFAS analysis was cleaned up by adding 20 μL acetic acid (and 20 mg ENVI Carb (Supelco) to the extract. The sample was vortexed for 30 s and centrifuged (3000 rpm) for 10 min. The extract was transferred to a new tube, evaporated until dryness, and reconstituted in 200 μL 50/50 MeOH/Milli-Q water (v/v).

The PFAS analysis in dust and serum extracts was performed using liquid chromatography coupled to tandem mass spectrometry (LC-MS/MS). The LC system was an Agilent 1200 Series (Palo Alto, CA, USA) coupled with an Agilent 6410 electro spray interface (ESI) operated in the negative ion mode prior to triple-quadrupole mass spectrometric detection. The details regarding LC and MS settings for cat serum (Table S3 and Table S4) and dust (Table S5 and Table S6) analysis are reported in detail in Additional file 1. The MS was operated in the electrospray negative ionization mode (ESI-) using two multiple reaction monitoring (MRM) transitions for each compound (Table S7 and Table S8). The total PFOS concentration was determined, i.e., the linear and the branched peaks were integrated together. The other branched isomers were not separated.

**QA/QC**

In parallel with the samples, solvent blank samples (n=10 for dust analysis and n=5 for cat blood analysis) were analyzed to determine background contamination. In addition, two serum sampling tubes were extracted and analyzed for possible leaching of PFASs from the material. To control for recovery, reproducibility, and accuracy, reference materials were included in the analysis, consisting of enriched (3 ng/mL) fetal bovine serum (50 μL, n=5, PAA laboratories, Pasching Austria) and NIST reference dust SRM2585 (50 mg, n=10). The results of the analytical quality assurance (QA) and quality control (QC) of the blood serum and dust sample analyses are reported in detail in Additional file 1 (Table S9-14 and Figure S1).

Average recoveries obtained in bovine serum enriched with PFSA, PFCA, 6:2 FTSA, and FOSA (at levels of 3 ng/mL) varied between 86 and 114% (relative SD <15%, n=8) depending on the PFAS (Table S9). For OP-PFASs, recoveries were determined in bovine serum enriched at two levels, 0.16 and 0.74 ng/mL. Three different tests to determine the influence of when to add the IS solution to the samples were performed and are reported in Additional file 1 (Figure S1). Recovery ranged from 50 for PFDP A to >200% for PFHxPA. Labeled IS recovery in the samples ranged between 32 and 259% (Table S11). Due to low recovery (<20%) of IS Cl-PFHxPA for PFPA and PFPIA quantification, these values are not reported.

The method limit of detection (LOD) was set to the lowest calibration concentration with a peak height larger than signal to noise of 3. The limit of quantification (LOQ) was set to 3 times the LOD or the average blank level + 3 times the standard deviation (SD) of the measured PFAS levels in solvent blank samples. The LOQs ranged between 0.2 and 170 pg/mL in blood serum (Table S10) and from 0.01 to 31 ng/g in dust (Table S14) depending on the specific PFAS.

The dust reference material SRM2585 was analyzed in replicates (n=8) and compared to PFAS levels reported in literature (Reiner et al. 2015). Accuracy was between 78% and 133% for PFSA and PFCA (Table S12). At the time, there was no literature data on the presence and levels of OP-PFASs in the SRM2585 reference material, and therefore, OP-PFASs were added to the material to evaluate the accuracy (between 3 and 3800 ng/g dust). Recovery of the OP-PFAS fortified samples ranged between 76 and 104% (Table S13). The standard deviation of all PFASs analyzed in the reference material was <20%, indicating a good reproducibility (Table S12 and S13).

### Statistical analysis

The data was checked for potential outliers using Tukey’s outer fence (Foreman 2014), but using 6 × IQR (interquartile range) instead of the original suggested 3 × IQR to get the filter more conservative. Despite this, several extreme values were found outside this range. To achieve a robust test, not influenced by these extremes, Spearman’s coefficient of rank correlation (rₚ) was used to correlate between serum PFAS concentrations (molar basis) and serum TH and cholesterol levels. For cases without obvious outliers, ordinary log-linear regression analysis and Pearson correlation (r) and the Coefficient of determination r², were applied. Corrections for ties were applied according to Zar 1999 (Zar 1999). p-values were calculated according to Press et al. (1996). Only positive correlations between serum and dust concentrations were assumed; therefore, one-tailed tests were applied. For the statistical evaluation, values >LOD but <LOQ the LOQ values (for that batch analysis) divided by square root of 2 were used. The non-parametric paired Wilcoxon signed-rank test (Wilcoxon 1945) was used to check for significant differences among rooms within the same homes. Multiple regression analysis (e.g., Kleinbaum et al. 2008) was used to study the potential effects of several PFASs on T4 levels. Principal component analyses (PCA e.g. Vannosa and Filzmoser 2009) were used to disclose potential patterns in relative PFAS concentrations among room types. A significance level of 5% (α = 0.05) was applied in the statistical analyses. The statistical analyses were carried out using the software package PIA (Bignert 2013).

### Results

Table 1 summarizes the PFAS concentrations determined in cat serum and household dust. In Additional file 1 (Table S15-25), the detailed information is reported on all individual blood serum and dust samples analyzed.
PFASs levels in cat serum

All samples (n=27) were analyzed for PFCAs, PFSAs, FOSA, and 6:2 FTSA levels, but two samples were lost during analysis of PFPAs, PFPIAs, and PAPs (n=25). Due to limited sample volume, no repeated analysis could be performed. PFOS, PFOA, PFDA, and PFUnDA could be detected in all cat blood samples. The highest median concentration was found for PFOS (2300 pg/mL), followed by PFOA (1100 pg/mL). Concentrations of PFPIAs and PAPs were an order of magnitude lower than PFSA/PFCA. The 6:2 and 8:2 diPAPs were detected in 68% (median 5.3 pg/mL) and 92% (median 8.7 pg/mL) of the samples, respectively. The 6:2 and 8:2 PAP concentrations could only be detected in 24% and 8% of the samples, respectively. Three perfluoroalkyl phosphinic acids (6:6-, 6:8-, and 8:8 PFPIA) could be determined in 12–28% of the cat serum samples. Unfortunately, the analytical recovery of the internal standard Cl-PFHxPA was too low (Table S11) during analysis of the cat serum to be able to quantify the PFPA. The levels of sum of all PFASs (ΣPFASs) in the cat serum ranged between 1600 and 23000 pg/mL.

PFASs levels in household dust

PFOA and PFOS were detected in the dust from all rooms and quantified in 93% of the rooms. Median PFOA level was 9 ng/g (max 650 ng/g), and median PFOS level was 13 ng/g (max 220 ng/g). PFNA and PFDA were detected in 100% of the dust samples from the living rooms, but to a lesser extent in

| PFASs levels in cat serum (n=27) and household dust (n=46) |
|---------------------------------|-------------------|-------------------|-------------------|-------------------|-------------------|-------------------|
| PFASs                          | >LOD (%) | >LOQ | Mean (pg/mL) | Median | Min-max | >LOD (%) | >LOQ | Mean (ng/g dust) | Median | Min-max |
| PFBA                           | 0        | 0    | -             | -      | -       | 11       | 0    | 2.0               | <LOQ   | <LOD-23 |
| PFPeA                          | 0        | 0    | -             | -      | -       | 7        | 0    | 0.93              | <LOQ   | <LOD-23 |
| PFHxA                          | 0        | 0    | -             | -      | -       | 96       | 52   | 13                | 6.4    | <LOD-99 |
| PFHpA                          | 96       | 78   | 1200          | 580    | <LOD-4300 | 57       | 24   | 5.3               | 2.2    | <LOD-95 |
| PFOA                           | 100      | 96   | 1800          | 1100   | <LOQ-15000 | 100      | 93   | 51                | 9.0    | <LOQ-650 |
| PFNA                           | 93       | 70   | 360           | 220    | <LOD-1000 | 93       | 54   | 6.2               | 3.4    | <LOD-44 |
| PFDA                           | 100      | 100  | 500           | 430    | 130-1500 | 96       | 52   | 10                | 3.2    | <LOD-100 |
| PFUnDA                         | 100      | 96   | 620           | 490    | <LOQ-1700 | 54       | 17   | 1.9               | 1.0    | <LOD-22 |
| PFDoDA                         | 96       | 70   | 140           | 120    | <LOD-290 | 41       | 24   | 6.5               | <LOQ   | <LOD-110 |
| PFTeDA                         | 0        | 0    | -             | -      | -       | 22       | 4    | 0.85              | <LOQ   | <LOD-9.1 |
| PFTrDA                         | 0        | 0    | -             | -      | -       | 30       | 17   | 4.1               | <LOQ   | <LOD-56 |
| PFBS                           | 0        | 0    | -             | -      | -       | 2        | 2    | 0.19              | <LOQ   | <LOD-8.7 |
| PFHxS                          | 96       | 96   | 530           | 460    | <LOD-2100 | 24       | 7    | 0.61              | <LOQ   | <LOD-8.1 |
| PFHpS                          | 41       | 22   | 36            | <LOQ   | <LOD-170 | 0        | 0    | -                 | -      | -      |
| PFOS                           | 100      | 100  | 2300          | 2300   | 680-5000 | 100      | 93   | 22                | 13     | <LOQ-220 |
| FOSA                           | 7        | 4    | 5.0           | <LOQ   | <LOD-100 | 0        | 0    | -                 | -      | -      |
| 6:2 FTSA                       | 0        | 0    | -             | -      | -       | 65       | 30   | 10                | 1.8    | <LOD-220 |
| 6:2 PAP                        | 24       | 16   | 15            | <LOQ   | <LOD-130 | 100      | 87   | 67                | 31     | <LOQ-460 |
| 8:2 PAP                        | 8        | 4    | 4.8           | <LOQ   | <LOD-75  | 100      | 93   | 47                | 22     | <LOQ-360 |
| 6:6 PFPIA                      | 16       | 16   | 0.5           | <LOQ   | <LOD-17  | 39       | 26   | 1.0               | <LOQ   | <LOD-16 |
| 6:8 PFPIA                      | 28       | 28   | 2.2           | <LOQ   | <LOD-5.2 | 48       | 33   | 1.7               | <LOQ   | <LOD-27 |
| 8:8 PFPIA                      | 12       | 12   | 0.12          | <LOQ   | <LOD-1.1 | 61       | 35   | 0.73              | 0.10   | <LOD-11 |
| 6:2 diPAP                      | 68       | 40   | 11            | 5.3    | <LOD-130 | 100      | 100  | 140               | 65     | 11-1300 |
| 8:2 diPAP                      | 92       | 48   | 13            | 8.7    | <LOD-62  | 100      | 100  | 100               | 49     | 7-920 |
| PFHxPA                         | n.d.     | -    | -             | -      | -       | 39       | 20   | 3.3               | <LOQ   | <LOD-31 |
| PFOPA                         | n.d.     | -    | -             | -      | -       | 70       | 41   | 47                | 3.3    | <LOD-1800 |
| PFDoPA                         | n.d.     | -    | -             | -      | -       | 43       | 28   | 8.7               | <LOQ   | <LOD-70 |
| Σ PFASs                        | 7600     | 6400 | 1600-23000    | 550    | 280     | 50-2400  |      |                  |        |        |

n.d. not determined
the other rooms (Table S18). 6:2 and 8:2 diPAPs were detected and quantified in all rooms from all homes, whereas the 6:2 and 8:2 PAPs were detected in all rooms and quantified in 87% and 93% of the rooms. The median concentrations for diPAPs (65 and 49 ng/g for 6:2 and 8:2 diPAP, respectively) are higher than for PFOA and PFOS, which are dominating the profile for PFCA and PFSA, respectively. PFOPA was detected in 70% of the dust from all rooms but only quantified in 41% of the samples with a median of 3.3 ng/g. There was a larger difference regarding OP-PFAS concentrations between

Fig. 1 $\sum$PFAS levels (ng/g) in the dust from different rooms sampled at the 17 participating families

Fig. 2 PFAS profile (median) in cat serum (pg/mL) and household dust (ng/g dust)
families than for PFCA and PFSA. One family had elevated levels in the living room, with a PFOPA level of 1800 ng/g dust (family 8, Table S 18, Fig. 1). The highest $\sum$PFASs level was found in the child’s bedroom of family 15, due to elevated diPAP levels (Table S23). Family 2 reported elevated levels of both PAPs and diPAPs in the living room (Table S 19).

Statistical significance was found when applying the Wilcoxon signed-rank test, which is avoiding variation among homes, hence increasing the possibility to detect potential differences among rooms within the same homes. The test showed that 8:8 PFPIA concentrations were higher in children’s rooms compared to adults’ and also that PFHxA and PFUnDA concentrations were lower (Figure S 2). Further, 8:2 PAP concentrations were indicated to be higher in living rooms compared to children’s rooms (Figure S 3), as well as 6:2 FTSA concentrations compared to adults’ rooms (Figure S4). Although the concentrations differed between rooms for PFASs, there was no significant difference regarding the PFAS profiles between rooms (Figure S5).

**PFASs levels in household dust vs. cat serum**

The levels measured in household dust and in cat blood serum were statistically tested for association between the two matrices, evaluating whether dust is an exposure pathway to cats. In general, the PFAS profile showed resemblances between the two matrices, although the relative concentrations of different PFAS groups differed (Fig. 2). The most significant difference was the dominance of PFOS and PFOA in cat serum and of PAPs and diPAPs in household dust.

Only 8 PFASs (PFHpA, PFOA, PFNA, PFDA, PFUnDA, PFOS, 6:2 diPAP, and 8:2 diPAP) were detected (>LOD) in more than 50% of both cat serum and house dust and were paired for comparison (Figure S6 and S7). As there was no difference between the rooms, the average PFAS dust level was used to compare with the cat(s) living in that household. Serum concentrations were not age-adjusted as no significant correlation could be found with age, although PFOS ($p=0.068$) and PFHxS ($p=0.090$) were close to being significant. All levels were compared using molar-based concentrations. Significant positive correlations were only found for PFOA ($r_s=0.32, p<0.049$) and PFUnDA ($r_s=0.55, p<0.001$), confirming that dust is a relevant PFAS exposure pathway for cats.

**PFASs levels in cat serum vs. thyroid hormone levels**

The cats represented a wide range of breed, age, and size (Norrgran Engdahl et al. 2017). The cats were all healthy, and measured thyroid hormone status (total T4 and TSH) was within a normal range (Table S15). A significant positive increase of serum total T4 levels could be observed with increasing PFNA levels (Fig. 3). The ratio between T4 and TSH levels is sometimes a more sensitive indication to changes in the thyroid hormone system due to feedback regulation, e.g., lowered T4 will increase the response in TSH and vice versa. It has been shown that PFASs bind to the T4 transport protein TTR (Weiss et al. 2009). Here we calculated the total T4 equivalency (T4 eq) by multiplying the T4 relative potency (T4 REP) determined in a radioligand binding assay (Weiss et al. 2009) by the individual PFAS concentrations ($C_i$) and sum it up for the individual serum samples (T4 eq = $\Sigma[C_i \times T4 REP_i]$). Both the serum PFAS and total T4 eq levels were compared to T4, TSH, or the ratio between the two, but no correlations could be seen (Figure S8).

**PFASs levels in cat serum vs. cholesterol levels**

EFSA recently established health-based guidance values for PFOS and PFOA, where one of the critical effects were the increase in serum total cholesterol (EFSA 2018). Mainly PFOS and PFOA have been studied regarding how blood lipid metabolism is affected by PFASs. Here we could not see a statistically significant increase of the cholesterol associated to PFOS and PFOA, although to the total PFASs levels (Fig. 4), as well as with PFHpA, PFUnDA, and PFDoDA (Figure S9). No correlations between cat serum triglycerides and PFASs (individual and sum) were found.

**Discussion**

PFAS levels in household dust were within the same range as previously reported. It can be a challenge to compare PFAS levels determined in dust from different studies, as levels are...
highly influenced by sampling design, e.g., how the dust is collected, the analyzed sample size fraction, region, as well as sampling year. Here we chose to compare our reported PFAS levels with a study where pooled household dust from 8 countries was sampled worldwide in the same sampling year as our study (2013–2014) and analyzed for a wide range of PFASs (Eriksson and Kärrman 2015). The particle size fraction was smaller (<150 μm) than in our study (40–1000 μm), which could influence the reported results. An increasing surface-to-area ratio with decreasing particle size can adsorb more analytes (Weiss et al. 2018). Australia, Faroe Islands, and Sweden showed similar PFAS profiles in dust. Except Japan and Canada, the other countries have no PFAS production sites, and the results indicate that consumer products used in the homes are significant sources of PFASs (Wang et al. 2014). Dust from Canada and Australia contained higher dust concentrations of PFOS and PFHxS, chemicals typically used as surfactants in firefighting foams, for example. PFNA was higher in Canada and Japan, a chemical used for the production of polyvinylidene fluoride (Wang et al. 2014). 6:2 and 8:2 diPAP dominated the PFAS profiles in dust from all countries, except Greece, Nepal, and Spain, which were the countries with the lowest total PFAS levels. PAPs were not included in the comparison as they were only semi-quantified in the study. The indicated median levels of PAPs (6:2, 8:2, and 10:2 PAP) in the different countries ranged from 3.7 to 1 023 ng/g, and the levels were dominating the PFAS profile together with diPAPs (Σ15 diPAPs, 3.6–692 ng/g). As diPAPs dominates the technical mixture used in products, it is likely that PAPs are products of degradation of diPAPs. (Lee et al. 2010).

Eight PFASs could be paired in the dust and cat serum (Figure S6). Two PFASs, PFOA, and PFUnDA were significantly correlated, demonstrating that dust was a relevant exposure pathway. It is reasonable to speculate that other exposure routes, such as drinking water and cat food, are contributing to cat serum levels, especially for the levels of PFOS and PFHxS commonly found in fish and drinking water (Ahrens et al. 2015). Therefore, the correlations found for PFOA and PFUnDA indicate that dust is an especially relevant exposure pathway for these compounds and that, generally, chemicals associated to dust are ingested.

Regarding PFCAs and PFSAs in cat serum, the levels are comparable to other studies reporting on PFASs in cats and humans (Fig. 6). There are only two other studies reporting on PFASs in cats, on cats situated in the USA (Bost et al. 2016; Wang et al. 2018). The levels were higher than in the Swedish cats and seem to be decreasing over time, although the number of samples was too small to draw any conclusions. PFASs were determined in children from Finland at different ages while growing up (Koponen et al. 2018). Levels were decreasing with age, which was explained by the rapid body weight gain whereas the total body load was constant. At the age of 10, the average serum level of PFOA was similar to the median cat serum levels in Sweden, and at the age of 6 for PFOS. Serum has been analyzed from 1st-time mothers coming from the same region as the cats in this study. PFAS levels were similar except for PFHxS and PFOS, which were equally high as in cats in the USA in 2008. This was due to drinking water contamination from a firefighting training site (Ahrens et al. 2015). The cats from this study coming from that specific region did not have elevated PFHxS and PFOS levels, probably due to the fact that cats drink very little water. Their main water supply comes via their food.

The dominance of PAPs and diPAPs in dust was not reflected in the cat serum. Only low levels of PAPs and diPAPs were determined in a limited number of serum samples. PFASs, including diPAPs, have been determined in human serum samples from donors in the USA in 2009 (Lee and Mabury 2011). The levels of PFOS and PFOA were comparable to the reported cat serum samples here, but diPAP levels were about 10 times higher and were found in 80% of the samples. This indicates that PAPs and PFSA/PFCA have different sources. PAPs were not determined in the US blood donors. The biotransformation of diPAPs has been studied in Carp (Chen et al. 2019). Several metabolites were detected but not the corresponding PAPs. It has been suggested that PAPs in a typical mammalian digestive tract would hydrolyze in approximately 100 s (Jackson and Mabury 2012). In an occupationally exposed population in China, average 6:2 and 8:2 diPAP levels of 0.19 and 0.17 ng/mL and 6:2 PAP level of 0.11 ng/mL have been reported. PFOA and PFOS levels were considerably higher than in the general population (average of 325 and 1064 ng/mL, respectively), and it is
reasonable to believe that the workers were exposed to PAPs and diPAPs in their occupation as well (Gao et al. 2018). Serum, plasma, and whole blood were collected from men and women \((n=61)\) living in the Oslo area, Norway, to determine the distribution between the matrices of these surface-active compounds (Poothong et al. 2017). It was observed that the quantification frequency was much higher in plasma than in serum, which is the matrix commonly analyzed from whole blood had the lowest quantification frequency for PAPs, clearly showing the complexity to extract these compounds from blood proteins.

It is worth noting that the cats in this study were all healthy, and therefore, any statistical association to altered thyroid hormone levels is challenging to find in a study with only a limited number of cats. Despite that, a positive significant association between the serum total T4 and PFNA levels

\[\text{Fig. 5} \quad \text{Dust PFAS levels (ng/g dust) compared between this study and one similar study (Eriksson and Kärrman 2015). Selected analytes have been determined in all samples, and the sampling years are 2013–2014. Asterisk shows the average concentration determined in pooled dust from vacuum cleaner consisting of “n” numbers of bags. The analyzed particle size fraction is <150 \(\mu\)m. § shows the arithmetic mean PFAS concentration of the dust sampled in this study \((n=46)\) and consist of still-standing dust of particle size 40–1000 \(\mu\)m.}\]

\[\text{Fig. 6} \quad \text{PFAS concentrations determined in cat serum and human serum taken from literature. (1) Geometric mean (Bost et al. 2016). (2) Geometric mean (Wang et al. 2018). (3) This study, median. (4) Average of median for male and female children (Koponen et al. 2018). (5) Pooled blood serum samples (pg/g serum) (Gebbink et al. 2015).}\]
was observed (Fig. 3). In children 1–17 years of age living in the vicinity of a chemical plant, this association was also observed (Lopez-Espinosa et al. 2012). Serum PFOA concentrations were also associated with thyroid disease in those children. In the two studies with cats from the USA, where different endocrine diseases were studied, a significant association between hyperthyroidism and PFOA levels was reported, as well as a weak association with $\sum$PFAS levels (Bost et al. 2016; Wang et al. 2018). No significant correlation could be determined between PFAS levels and the amount of time the cat spent indoors, as was observed in another study (Bost et al. 2016).

A recent study evaluated dust-related thyroid hormone disrupting compounds in mixtures corresponding to household dust, infant blood serum, and adult blood serum (Hamers et al. 2020). In total, 25 chemicals were measured for their binding potency to the transport protein TTR in a competitive in vitro assay (FITC-T4). An effect level corresponding to the inhibition of the T4-TTR binding of 20% was used in the study, mainly as robust data could be achieved at that level, but also as even small changes in available TH levels can cause severe effects on brain development. Moreover, animal studies with OHCs have reported an association between decreased maternal levels of circulating T4 and neurodevelopmental effects on cognitive function, motor activity, and behavior in offspring (EFSA 2005; EFSA 2011a; EFSA 2011b; EFSA 2018). By applying the toxic unit (TU) summation, which assumes the principle of concentration addition for the investigated 25 chemicals, it was predicted that from 1.3 up to 6.2% of T4-TTR binding in human blood could be inhibited by the chemical mixture tested (Hamers et al. 2020). The calculations took into consideration serum TBG and ALB levels as relevant TH binders in humans, where a 12/71/17% distribution of T4 over TTR/TBG/ALB was calculated. As TBG has not been found in cat serum, it is believed that TTR has a more important role in the transport of TH in cats than in other mammals (Larsson et al. 1985). This study is part of a project abbreviated MiSSE (Mixture Assessment of EDCs), and the same cat serum samples have been measured for a range of OHCs, summarized in Table 2. The TUs are calculated by dividing the measured concentration (median or max) by the effect concentration (IC20). If the sum of individual TUs equals 1, the concentration is expected to produce the effect (20% inhibition of T4-TTR binding).

The $\sum$TUs for TU$_{\text{median}}$ in the cat serum was 0.17, which is a margin of 5 from equal to 1, whereas the $\sum$TUs for TU$_{\text{max}}$ was 0.79. Just as in human blood (Hamers et al. 2020), PFOS was one of the major contributors to the $\sum$TUs in cat serum. The PFASs in cat serum constituted almost 50% of the $\sum$TUs for the median levels, and 36% for the max values, despite the rather low binding potency of PFASs (Weiss et al. 2009). It is difficult to extrapolate the in vitro assay results to real-life conditions and to estimate the relevance of this T4 displacement from TTR in cats remains to be clarified.

This exercise only contains 13 compounds determined in the cat’s serum of this project. In the literature, more than 60 OHCs have been reported in cat serum, of which many have been tested for their binding potential to TTR (Weiss and Jones 2020). It is therefore likely to believe that the TU summation here is an underestimation of the real situation for household cats. Only considering the PFASs on the market, PFAS composition is changing in products as regulations change and new PFAS substitutes are entering the market. Today there are believed to be more than 3000 different PFASs circulating the global market, and more than 4500

| Substance | IC$_{20}$ (nM) | $C_{\text{median}}$ (nM) | $C_{\text{max}}$ (nM) | TU$_{\text{median}}$ | TU$_{\text{max}}$ |
|-----------|----------------|--------------------------|------------------------|----------------------|------------------|
| PFHxS     | 136            | 1.2                      | 5.3                    | 0.01                 | 0.04             |
| PFOS      | 82             | 4.6                      | 10                     | 0.06                 | 0.12             |
| PFHpA     | 299            | 1.6                      | 12                     | 0.01                 | 0.04             |
| PFOA      | 491            | 2.8                      | 35                     | 0.01                 | 0.07             |
| PFNA      | 361            | 0.47                     | 2.2                    | 0.001                | 0.01             |
| PFDA      | 1710           | 0.84                     | 2.9                    | 0.0005               | 0.002            |
| PFUnDA    | 1010           | 0.87                     | 3.0                    | 0.0009               | 0.003            |
| Pentachlorophenol $^b$ | 17        | 0.41                     | 4.2                    | 0.02                 | 0.24             |
| CB153 $^b$ | 77             | 0.17                     | 1.5                    | 0.002                | 0.02             |
| 2,4,6-TBP $^b$ | 8.9        | 0.44                     | 1.1                    | 0.05                 | 0.13             |
| 6OH-BDE47 $^b$ | 31        | 0.5                      | 3.4                    | 0.02                 | 0.11             |
| BDE47 $^b$ | 210            | 0.04                     | 0.26                   | 0.0002               | 0.001            |
| BDE99 $^b$ | 73             | 0.06                     | 0.61                   | 0.0008               | 0.008            |
| $\sum$TUs |                |                          |                        | 0.17                 | 0.79             |

$^a$ Hamers et al. (2020)  
$^b$ Norrgran Engdahl et al. (2017)
PFASs are registered with CAS numbers (KemI 2015; OECD 2018; Wang et al. 2017). What yet needs to be evaluated is whether the new PFASs are bioavailable and could pose a threat to health.

Several epidemiological studies have reported associations between PFASs (mainly PFOS and PFOA) and human health outcomes linked to blood lipids (Eriksen et al. 2013; Fletcher et al. 2013; Li et al. 2020; Liu et al. 2020). The underlying mechanisms are discussed, but it is agreed on that interferences with the lipid metabolism may potentially lead to an elevated risk of developing cardiovascular diseases, which is one of the 6 most common diagnostic categories for mortalities in cats (Egenvall et al. 2009). It is out of the scope for this study to speculate why some PFASs are correlating with the serum cholesterol and some are not. It is important to acknowledge that the sample size in this study is limited and hence the results should be understood as indicative and further research is needed before prompt conclusions can be drawn.

Conclusion

This study confirmed that dust is a relevant exposure pathway to PFASs for cats. Levels are similar in humans and pet cats, and it is plausible that the exposure to certain PFASs can affect the thyroid hormone system, although that needs to be confirmed with further research. The critical health effect is recognized by EFSA; increased total cholesterol in blood could also be observed in cats, although for total PFAS, PFHpA, PFUnDA, and PFDoDA, and not for PFOS and PFOA. The high levels of PAPs and diPAPs in dust samples were not reflected in the cats’ serum, indicating a lower bioavailability, fast excretion, or metabolism to PFCAs. With respect to PFASs, the potential of using cats as a model organism for human exposure has been demonstrated. By combining exposure assessment with health parameters in studying pet animals, we can improve our understanding of the health impacts that exposure to indoor-related chemicals have on us and our pets.

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Author contribution JW and BJ performed the sampling of the cat serum and household dust. JW is also the project leader of MiSSE, designed the study, and was leading in the writing of the manuscript. JK performed the chemical analyses under supervision of JW and ML. AB performed all statistical analysis. All authors contributed to and approved the final manuscript.

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Data availability All data generated or analyzed during this study are included in this published article and the supporting information in Additional file 1.

Declarations

Ethics approval and consent to participate The study was performed after permission from the Swedish Board of Agriculture and Uppsala Ethical Committee on Animal Studies (No. 31-10466/12). The participating families were volunteers and all provided a written consent.

Consent for publication Not applicable.

Competing interests The authors declare no competing interests.

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References

Ahrens L, Norström K, Viktoria T, Cousins AP, Josefsson S (2015) Stockholm Arlanda Airport as a source of per- and polyfluoroalkyl substances to water, sediment and fish Chemosphere 129:33-38. https://doi.org/10.1016/j.chemosphere.2014.03.136

Ali N, Malik RN, Mehdi T, Eqani SAMAS, Javeed A, Neels H, Covaci A (2013) Organohalogenated contaminants (OHCs) in the serum and hair of pet cats and dogs: Biosentinels of indoor pollution. Sci Total Environ 449:29–36. https://doi.org/10.1016/j.scitotenv.2013.01.037

Beesoon S, Genuis SJ, Benskin JP, Martin JW (2012) Exceptionally high serum concentrations of perfluorohexanesulfonate in a Canadian family are linked to home carpet treatment applications. Environ Sci Technol 46:12960–12967

Abbreviations OHC, Organohalogen compound; EDC, Endocrine disrupting compound; TH, Thyroid hormone; T4, Thyroxine; TSH, Thyroid stimulating hormone; TTR, Transthyretin; PFAS, Per- and polyfluoroalkyl substance; PFCA, Perfluoroalkyl carboxylic acid; PFSA, Perfluoroalkyl sulfonic acid; OP-PFAS, Organophosphorus PFAS; PFPA, Perfluoroalkyl phosphonic acid; PFPIA, Perfluoroalkyl phosphinic acid; PAP and diPAP, Polyfluoroalkyl phosphate mono- and diesters; FTSA, Fluorotelomer sulfonic acid; FOSA, Perfluoroctane sulfonamide; LOD, Limit of detection; LOQ, Limit of quantification; SPE, Solid-phase extraction; TU, Toxic unit

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Bignert A (2013) The PIA application for statistical analysis of time-series data. AMAP
Bost PC, Strynar MJ, Reiner JL, Zweigenbaum JA, Secoura PL, Lindstrom AB, Dye JA (2016) U.S. domestic cats as sentinels for perfluoroalkyl substances: possible linkages with housing, obesity, and disease. Environ Res 151:145–153. https://doi.org/10.1016/j.envres.2016.07.027
Buck RC et al (2011) Perfluoroalkyl and Polyfluoroalkyl substances in the environment: terminology, classification, and origins. Integr Environ Assess Manag 7:513–541. https://doi.org/10.1002/ieam.258
Chen M et al (2019) Biotransformation and bioconcentration of 6:2 and 8:2 perfluorooalkyl phosphate diesters in common carp (Cyprinus carpio): underestimated ecological risks. Sci Total Environ 656:201–208. https://doi.org/10.1016/j.scitotenv.2018.11.297
Chow K, Hearn LK, Zuber M, Beatty JA, Mueller JF, Barrs VR (2015) EFSA (2011a) Scientific Opinion on Polybrominated Diphenyl Ethers (PBDEs). EFSA J 9:1
Covaci A, Voorspoels S, Thomsen C, Van Bavel B, Neels H (2006) Evaluation of total lipids using enzymatic methods for the normalization of persistent organic pollutant levels in serum. Sci Total Environ 366:361–366
Dirtu AC, Niessen SJM, Jorens PG, Covaci A (2013) Organohalogenated contaminants in domestic cats’ plasma in relation to spontaneous acromegaly and type 2 diabetes mellitus: a clue for endocrine disruption in humans? Environ Int 57–58:60–67. https://doi.org/10.1016/j.envint.2013.04.004
Dye J, Venier M, Zhu L, Ward CR, Hites RA, Birnbaum L (2007) Elevated PBDE levels in pet cats: Sentinels for humans. Environ Sci Technol 41:6350–6356
EFSA (2005) Opinion of the scientific panel on contaminants in the food chain on a request from the commission related to the presence of non dioxin-like polychlorinated benzenes (PCB) in feed and food. EFSA J 1:1–137
EFSA (2011a) Scientific Opinion on Polybrominated Diphenyl Ethers (PBDEs) in Food. EFSA J 9:1–274. https://doi.org/10.2903/j.efsa.2011.2156
EFSA (2011b) Scientific opinion on tetrabromobisphenol A (TBBPA) and its derivatives in food. EFSA J 9:2477. https://doi.org/10.2903/j.efsa.2011.2477
EFSA (2018) Risk to human health related to the presence of perfluorooctane sulfonic acid and perfluorooctanoic acid in food. EFSA J 16:5194. https://doi.org/10.2903/j.efsa.2018.5194
Egevall A, Nødtvedt A, Häggström J, Ström Holst B, Möller L, Bonnett BN (2009) Mortality of life-insured Swedish cats during 1999-2006: age, breed, sex, and diagnosis. J Vet Intern Med 23:1175–1183. https://doi.org/10.1111/j.1939-1676.2009.0039.d
Eriksen KT, Raaschou-Nielsen O, McLaughlin JK, Lipworth L, Tjønneland A, Overvad K, Sørensen M (2013) Association between plasma PFOA and PFOS levels and total cholesterol in a Middle-Aged Danish Population. PLoS One 8:e56969. https://doi.org/10.1371/journal.pone.0056969
Eriksson U, Kärnman A (2015) World-wide indoor exposure to perfluorooalkyl phosphate esters (PAPs) and other PFASs in household. Dust Environ Sci Technol 49:14503–14511
Fletcher T, Galloway TS, Melzer D, Holcroft P, Cipelli R, Pilling LC, Mondal D, Luster M, Harries LW (2013) Associations between PFOA, PFOS and changes in the expression of genes involved in cholesterol metabolism in humans. Environ Int 57–58:2–10. https://doi.org/10.1016/j.envint.2013.03.008
Foreman JW (2014) Data smart: using data science to transform information into insight. John Wiley & Sons
Gao K et al (2018) An integrated method for simultaneously determining 10 classes of per- and polyfluoroalkyl substances in one drop of human serum. Anal Chim Acta 999:76–86. https://doi.org/10.1016/j.aca.2017.10.038
Gebbink WA, Glynn A, Berger U (2015) Temporal changes (1997–2012) of perfluoroalkyl acids and selected precursors (including isomers) in Swedish human serum. Environ Pollut 199:166–173. https://doi.org/10.1016/j.envpol.2015.01.024
Guo W, Gardner S, Yen S, Petreas M, Park J-S (2016) Temporal changes of PBDE Levels in California house cats and a link to cat hyperthyroidism. Environ Sci Technol 50:1510–1518. https://doi.org/10.1021/acs.est.5b04252
Hamers T, Kortenkamp A, Scholze M, Molenaar D, Cenijn PH, Weiss JM (2020) Thrashyretin-binding activity of complex mixtures representing the composition of thyroid-hormone disrupting contaminants in house-dust and human serum. Environ Health Perspect 128:017015–017011. https://doi.org/10.1289/EHP5911
Henríquez-Hernández LA, Carreton E, Camacho M, Montoya-Alonso JA, Boada LD, Bernal Martín V, Falcón Cordon Y, Falcón Cordon S, Zumbado M, Lizardo OP (2017) Potential Role of Pet Cats As a Sentinel Species for Human Exposure to Flame Retardants. Front Vet Sci 4:79. https://doi.org/10.3389/fvets.2017.00079
Jackson DA, Mabury SA (2012) Enzymatic kinetic parameters for perfluorinated alkyl phosphate hydrolysis by alkaline phosphatase. Environ Toxicol Chem 31:1966–1971
Jones B, Norrgran Engdahl J, Weiss JM (2019) Are persistent organic pollutants important in the etiology of feline hyperthyroidism? A review. Acta Vet Scand 61:45. https://doi.org/10.1186/s13028-019-0478-9
Jones-Ortazo HA et al (2005) Is house dust the missing exposure pathway for PBDEs? An analysis of the urban fate and human exposures to PBDEs. Environ Sci Technol 39:5121–5130
Keml (2015) Förekomst och användning av högfluorerade ämnen och alternativ vol Report 6/15
Kleinbaum DG, Kupper LL, Muller KE (2008) Applied Regression Analysis and Other Multivariate methods. 4th edn. Brooks/Cole Cengage learning
Koponen J et al (2018) Longitudinal trends of per- and polyfluoroalkyl substances in children’s serum. Environ Int 129:519–599. https://doi.org/10.1016/j.envint.2018.09.006
Larsson M, Pettersson T, Carlstrom A (1985) Thyroid hormone binding in serum of 15 vertebrate species: isolation of thyroxine-binding globulin and prealbumin analogs. Gen Comp Endocrinol 58:360–375
Lee H, Mabury SA (2011) A pilot survey of legacy and current commercial fluorinated chemicals in human sera from United States donors in 2009. Environ Sci Technol 45:8067–8074. https://doi.org/10.1021/es200167q
Lee H, D’eon J, Mabury SA (2010) Biodegradation of polyfluoroalkyl phosphates as a source of perfluorinated acids to the environment. Environ Sci Technol 44:3305–3310. https://doi.org/10.1021/es9028183
Li Y, Barregard L, Xu Y, Scott K, Pineda D, Lindh CH, Jakobsson K, Fletcher T (2020) Associations between perfluorooalkyl substances and serum lipids in a Swedish adult population with contaminated drinking water. Environ Health 19:33. https://doi.org/10.1186/s12940-020-05588-9
Liu G, Zhang B, Hu Y, Rood J, Liang L, Qi L, Bray GA, DeJonge L, Liu Z, Pineda D, Lindh CH, Jakobsson K, Mabury SA (2010) Biodegradation of polyfluoroalkyl phosphates as a source of perfluorinated acids to the environment. Environ Sci Technol 44:3305–3310. https://doi.org/10.1021/es9028183
Lopez-Espinosa M-J, Mondal D, Armstrong B, Bloom MS, Fletcher T (2012) Thyroid function and perfluoroalkyl acids in children living near a chemical plant. Environ Health Perspect 120:1036–1041. https://doi.org/10.1289/ehp.1104370
Mizukawa H, Nomiyama K, Nakatsu S, Iwata H, Yoo J, Kubota A, Yamamoto M, Ishizuka M, Ikenaka Y, Nakayama SMM, Kunisue T, Tanabe S (2016) Organohalogen Compounds in pet dog and cat: do pets biotransform natural brominated products in food to harmful hydroxlated substances? Environ Sci Technol 50:444–452. https://doi.org/10.1021/acs.est.5b04216

Mizukawa H et al. (2017) Anthropogenic and naturally produced brominated phenols in pet blood and pet food in Japan. Environ Sci Technol 51:11354–11362. https://doi.org/10.1021/acs.est.7b01009

Norrgran Engdahl J, Bignert A, Athanassiadis I, Bergman Å, Weiss JM (2017) Cats’ Internal exposure to selected brominated flame retardants and organochlorines correlated to house dust and cat food. Environ Sci Technol 51:3012–3020. https://doi.org/10.1021/acs.est.6b05025

Norrgran J, Jones B, Bignert A, Athanassiadis I, Bergman Å (2015) Higher PBDE serum concentrations may be associated with feline hyperthyroidism in Swedish cats. Environ Sci Technol 49:5107–5114. https://doi.org/10.1021/acs.est.5b00234

OECD (2018) Toward a new comprehensive global database of per- and polyfluoroalkyl substances (PFASs): Summary report on updating the OECD 2007 list of PFASs. Paris

Peterson ME, Ward CR (2007) Etiopathologic findings of hyperthyroidism in cats. Vet Clin Small Anim 37:633–645

Poothong S, Thomsen C, Padilla-Sanchez JA, Papadopoulou E, Haug LS (2017) Distribution of novel and well-known poly- and perfluoroalkyl substances (PFASs) in human serum, plasma, and whole blood. Environ Sci Technol 51:13388–13396. https://doi.org/10.1021/acs.est.7b03299

Press WH, Teukolsky SA, Vetterling WT, Flannery BP (1996) Numerical recipes in FORTRAN 77. The Art of Scientific Computing. 2nd edn. Cambridge University Press

Reiner JL et al (2015) Polyfluorinated substances in abiotic standard reference materials. Anal Bioanal Chem 407:2975–2983. https://doi.org/10.1007/s00216-013-7330-2

Richardson SJ (2009) Evolutionary changes to transthyretin: evolution of transthyretin biosynthesis. FEBS J 276:5342–5356

Schreiber G, Southwell BR, Richardson SJ (1995) Hormone delivery systems to the brain - transthyretin. Exp Clin Endocrinol Diabetes 103:75–80

US-EPA (2017) Child-specific Exposure Factors Handbook vol Report No: EPA/600/R-17/384F. Environmental Protection Agency, National Center for Environmental Assessment, Washington DC, U.S.

Varnosa K, Filzmoser P (2009) Introduction to multivariate statistical analysis in chemometrics. Taylor & Francis

Vestergren R, Cousins IT (2009) Tracking the pathways of human exposure to perfluorocarboxylates. Environ Sci Technol 43:5565–5575. https://doi.org/10.1021/es900228k

Walter KM, Lin Y-P, Kass PH, Puschner B (2017) Association of polybrominated diphenyl ethers (PBDEs) and polychlorinated biphenyls (PCBs) with hyperthyroidism in domestic felines, sentinel for thyroid hormone disruption. BMC Vet Res 13:120. https://doi.org/10.1186/s12917-017-1031-6

Wang Z, Cousins IT, Scheringer M, Buck RC, Hungerbühler K (2014) Global emission inventories for C4–C14 perfluoroalkyl carboxylic acid (PFCA) homologues from 1951 to 2030, Part I: production and emissions from quantifiable sources. Environ Int 70:62–75. https://doi.org/10.1016/j.envint.2014.04.013

Wang Z, DeWitt JC, Higgins CP, Cousins IT (2017) A Never-Ending Story of Per- and Polyfluoroalkyl Substances (PFASs)? Environ Sci Technol 51:2508–2518. https://doi.org/10.1021/acs.est.6b04806

Wang M, Guo W, Gardner S, Petreas M, Park JS (2018) Per- and polyfluoroalkyl substances in northern California cats: temporal comparison and a possible link to cat hyperthyroidism. Environ Toxicol Chem 37:2523–2529

Weiss JM, Jones B (2020) Using cats as sentinels for human indoor exposure to organic contaminants and potential effects on the thyroid hormone system. In: Ramiro Pastorinho MaS ACA (ed) Pets as Sentinels, Forecasters and Promoters of Human Health. Springer Nature, Switzerland, pp 123–139

Weiss JM, Andersson PL, Lamoree MH, Leonards PE, van Leeuwen SP, Hamers T (2009) Competitive binding of poly- and perfluorinated compounds to the thyroid hormone transport protein transthyretin. Toxicol Sci 109:206–216. https://doi.org/10.1093/toxsci/kfp055

Weiss JM, Andersson PL, Zhang J, Simon E, Leonards PEG, Hamers T, Lamoree MH (2015) Tracing thyroid hormone disrupting compounds: database compilation and structure activity evaluation for an effect-directed analysis of sediment. Anal Bioanal Chem 407:5625–5634

Weiss JM, Gustafsson Å, Gerde P, Bergman Å, Lindh CH, Krais AM (2018) Daily intake of phthalates, MEHP, and DINCH by ingestion and inhalation. Chemosphere 208:40–49. https://doi.org/10.1016/j.chemosphere.2018.05.094

White JT, Kelly JW (2001) Support for the multigenic hypothesis of amyloidosis: the binding stoichiometry of retinol-binding protein, vitamin A, and thyroid hormone influences transthyretin amyloidogenicity in vitro. Proc Natl Acad Sci U S A 98:13019–13024

WHO/UNEP (2013) State of the science of endocrine disrupting chemicals - 2012. Geneva, Switzerland

Wilcoxon F (1945) Individual comparisons by ranking methods. Biom Bull 1:80–83

Wu X et al. (2015) Serum concentrations of perfluorinated compounds (PFC) among selected populations of children and Adults in California. Environ Res 136:264–273. https://doi.org/10.1016/j.envres.2014.09.026

Zar JH (1999) Biostatistical analysis, 4th edn. Prentice-Hall, Englewood Cliffs

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