Review RoTS analyses

Introduction

This notebook goes into several issues with the RoTS, 2018 memo on deriving a norm for PFOS contamination in soil at the construction site of Oosterweel.

The overall logic to determine a soil PFOS norm is

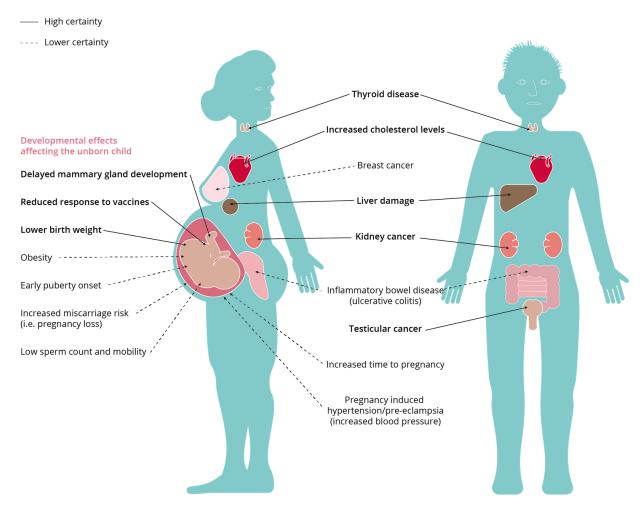
- Determine a safe level of todal daily intake (TDI) of PFOS for humans
- Attribute the maximal amount of PFOS intake through drinking water given all other sources of PFOS intake (primarily food)
- Next, calculate a maximal concentration of PFOS in drinking water based on an assumed 2.2L drinking per day (WHO standard)
- To guarantee the use of groundwater for drinking purposes, equate the maximal PFOS concentration in groundwater to the drinking water norm
- Based on experiments and literature data: compute the maximal PFOS in soil to guarantee leaching concentration at or below the drinking water norm

Health impacts of PFOS

The state of the art of scientific knowledge on the health impacts of PFOS and other PFAS is summarized in the abstract of the critical review from Fenton, et al., 2021 (image source: https://www.eea.europa.eu/public ations/emerging-chemical-risks-in-europe/emerging-chemical-risks-in-europe):

Reports of environmental and human health impacts of per-and polyfluoroalkyl substances (PFAS) have greatly increased in the peer-reviewed literature. The goals of the present review are to assess the state of the science regarding toxicological effects of PFAS and to develop strategies for advancing knowledge on the health effects of this large family of chemicals. Currently, much of the toxicity data available for PFAS are for a handful of chemicals, primarily legacy PFAS suchas perfluorooctanoic acid and perfluorooctane sulfonate. Epidemiological studies have revealed associations between ex-posure to specific PFAS and a variety of health effects, including altered immune and thyroid function, liver disease, lipid andinsulin dysregulation, kidney disease, adverse reproductive and developmental outcomes, and cancer. Concordance with experimental animal data exists for many of these effects. However, information on modes of action and adverse outcome pathways must be expanded, and profound differences in PFAS toxicokinetic properties must be considered in understanding differences in responses between the sexes and among species and life stages. With many health effects noted for a relatively few example compounds and hundreds of other PFAS in commerce lacking toxicity data, more contemporary and high-throughput approaches such as read-across, molecular dynamics, and protein modeling are proposed to accelerate the development of toxicity information on emerging and legacy PFAS, individually and as mixtures. In addition, an appropriate degree of precaution, given what is already known from the PFAS examples noted, may be needed to protect human health. (Fenton, et al., 2021)

Figure 1. Effects of PFAS on human health



 $Figure\ source:\ https://www.eea.europa.eu/publications/emerging-chemical-risks-in-europe/emerging-chemical-risks-in-europe$

Todal daily intake (TDI) for PFOS

Based on increasingly abundant evidence of adverse health impacts of PFOS and other PFAS, several regulating bodies have been ever more restrictive in their advises. Below we list the todal daily intake (TDI) values put out over the last decade by the Environmental Protection Agency (EPA, US) and the European Food Security Agency (EFSA, EU). Note that the adverse effects of PFOS on the immune system is long known in mice, and has more recently been corroborated by cohort studies during vaccination programs. It is exactly based on the latter that the latest more stringent norm of EFSA is based.

source	year	TDI (PFOS)	unit
EFSA	2008	150	ng/kg lg/dag
EPA	2009	80	ng/kg lg/dag
EPA	2016	20	ng/kg lg/dag
EFSA	2018	20	ng/kg lg/dag
EFSA	2020	0.63	ng/kg lg/dag

Attribution

As for most toxic contaminants, humans are exposed to PFAS via different pathways (food, water, dust, skin,...). De Silva et al. (2021) provide a critical review of the current knowledge on exposure pathways. In many cases, the exposure to or in take of chemical contaminants from drinking-water is much lower than that from other sources (WHO, 2011). Consequently, when deriving health norms for contaminants in certain products or media, one has to take the other exposure pathways into account. This is called attribution. The WHO has clear guidelines for how to determine attribution factors for contaminants in drinking water.

Abscence of exposure data

In the absence of adequate exposure data, the normal allocation of the total daily intake to drinking-water is 20%, which reflects a reasonable level of exposure based on broad experience, while still being protective. [...] There is variation in both the volume of water consumed daily and the body weight of consumers. It is therefore necessary to apply some assumptions in order to determine a guideline value. The default assumption for consumption by an adult is 2 litres of water per day, whereas the default assumption for body weight is 60 kg. In some cases, the guideline value is based on children, where they are considered to be particularly vulnerable to a particular substance. In this event, a default intake of 1 litre is assumed for a body weight of 10 kg; where the most vulnerable group is considered to be bottle-fed infants, an intake of 0.75 litre is assumed for a body weight of 5 kg. (WHO, 2011, p163-p164)

This heuristics is the standard procedure to compute the concentration of contaminants in drinking water in the abscense of exposure data. Simply by multiplying the TDI with body-weight and default attribution factor (20%), and dividing by a typical value of 2L per day for adults and 1L per day by children, we can expand the former table. Note that the final concentrations in drinking water are in nanogram per liter.

COURGO	voor	TDI (PFOS)	unit	drinking water, adult (60kg)	drinking water, child (10kg)	unit
source	year	(1105)	umi	drinking water, adult (ookg)	(10kg)	uiii t
EFSA	2008	150	ng/kg lg/dag	900	300	ng/L
EPA	2009	80	ng/kg lg/dag	480	160	ng/L
EPA	2016	20	ng/kg lg/dag	120	40	ng/L
EFSA	2018	20	ng/kg lg/dag	120	40	ng/L
EFSA	2020	0.63	$\frac{\mathrm{ng/kg}}{\mathrm{lg/dag}}$	3.78	1.26	ng/L

Exposure data

Obviously:

Wherever possible, data on the proportion of total daily intake normally ingested in drinking-water (based on mean levels in food, drinking-water and air) or intakes estimated on the basis of physical and chemical properties of the substances of concern are used in the derivation of guideline values. (WHO, 2011, p163-p164)

Since exposure pathway data exists for PFOS, we can do better than the heuristics above. We refere to 2 sources of exposure data: the review by Da Silva, et al (2021), and the EFSA (2020) report.

Table 2 of the Da Silva et al. (2021) is reproduced below. Outside Northern America (where relative contributions of 7 and 22% have been reported), the reported exposure via drinking water is very low, from less that 1% to 4%. (Note that, when reported, the intake via dust is of the same order of magnitude as the

intake via drinking water).

TABLE 2: Literature estimates of source contributions (percentage) to adult exposures to poly- and perfluoroalkyl substances^a

			E>	posure me	edium ^b	Exp	Exposure route ^b			
PFAS	Carbon length	Diet	Dust	Water	Consumer goods	Inhalation	Dermal	Indirect	Study location	Ref.
PFBA	4		4	96					NA	С
PFHxA	6	38	4	38		8		12	NA	С
PFHxA	6	87	4			2			Norway	d
PFHxS	6	57	38			5			Finland	e
PFHxS	6	94	1						Norway	d
PFHpA	7	93	1						Norway	d
PFH _p S	7				100				Norway	d
PFOA	8	16	11		58	14			NA, EŬ	f
PFOA	8	85	6	1	3			4	Germany, Japan	g
PFOA	8	77	8	11		4			Norway	h
PFOA	8	66	9	24		<1	<1		USA	i
PFOA	8	41		37				22	Korea	j
PFOA	8	99		<1					China	k
PFOA	8	47	8	12		6		27	NA	С
PFOA	8	95	<2.5			<2.5			Finland	е
PFOA	8	89	3			2			Norway	d
PFOA	8	91		3		5			Ireland	- 1
PFOS	8	66	10	7		2		16	NA	С
PFOS	8	72	6	22		<1	<1		USA	m
PFOS	8	96	1	1		2			Norway	h
PFOS	8	81	15		4				NA, EÚ	f
PFOS	8	93		4				3	Korea	j
PFOS	8	100		<1					China	k
PFOS	8	95	<2.5			<2.5			Finland	е
PFOS	8	75				3			Norway	d
PFOS	8	100							Ireland	- 1
PFOPA	8		100						Norway	d
PFNA	9	79	5			1			Norway	d
PFDA	10	51	2	4		15		28	NA	С
PFDA	10	78	1			2			Norway	d
PFDS	10		89		4				Norway	d
PFUnDA	11	61	4			1			Norway	d
PFDoDA	12	86	2	2		4		5	NA	С
PFDoDA	12	48	15						Norway	d
PFTrDA	13	89	1						Norway	d

Source: Da Silva, 2021

The relative low contribution of PFOS exposure via drinking water is confirmed in the EFSA, 2020 report. Based on data provided by the EU member states, the EFSA compiled PFOS exposure via food and drinking water. As can be seen on the compilation figure below, the intake of PFOS via drinking water is very small compared to other intake pathways, particularly compared to intake via seafood, fish and eggs. Based on the data provided by the EFSA (EFSA, 2020 - data appendix), we calculate that **the exposure via drinking water is only about 1%**, **both for adults as for toddlers, and also both on average in the EU as in Belgium.** Note that this number even neglects the intake via dust.

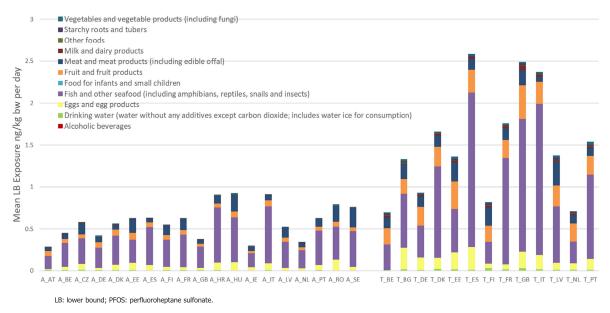


Figure 8: Mean LB exposure from different food categories of adults (A, left) and toddlers (T, right) to PFOS, for various surveys

Source: EFSA, 2020

Such low attribution factors are not special, but correspond to WHO guidelines for certain pesticides: > In the case of some pesticides, which are likely to be found as residues in food from which there will be significant exposure, the allocation for water may be as low as 1%. (WHO, 2011 p164)

But this has important ramifications for deriving a drinking water quality standard. Using the evidence based attribution factor of 1%, instead of the default value of 20%, we find the following drinking water concentrations:

source	year	TDI (PFOS)	unit	drinking water, adult (60kg)	drinking water, child (10kg)	unit
EFSA	2008	150	ng/kg lg/dag	45	15	$\overline{\mathrm{ng}/\mathrm{L}}$
EPA	2009	80	ng/kg lg/dag	24	8	ng/L
EPA	2016	20	ng/kg lg/dag	6	2	ng/L
EFSA	2018	20	ng/kg lg/dag	6	2	ng/L
EFSA	2020	0.63	$\frac{g}{g}$	0.189	0.063	ng/L

Partitioning and leaching from contaminated soil

Literature study

To guarantee the use of groundwater for drinking purposes, the maximal PFOS concentration in groundwater is set equal to the maximal concentration in drinking water. Subsequently, soils that can be re-used without treatment or special containment measures, we require that concentration in leaching water is also equal or below the drinking water standard. To determine the corresponding total contamination in the (in ng per kg dry weight) we need to know how much of the contaminant sticks to the soil (adsorption) and how much is freely available in the pore water. The so-called partitioning factors (Kd), which are nothing more

than the ratio's between soil concentration (C_s) and pore water (C_w) concentration of a contaminant under equilibrium (K_d = c_s/c_w). With those factor we can compute the relative fraction of adsorbed contaminant and contaminant in solution; they are determined experimentally either in the lab or in the field.

For many contaminants, partitioning is dependent on the soil properties (organic carbon (OC) content, pH, clay content, etc). This is also the case for PFOS. Li et al (2018) provide a critical review of the literature on partitioning of PFOS and other PFASs. In their Figure 1 they plot observed Kd values against soil organic matter content. Although there is an overall increasing trend of Kd with increasing organic matter content (they report a relatively high R^2 of 0.75 for the regression of PFOS vs. OC content) it can also be seen that there is a lot of variability in the reported values.

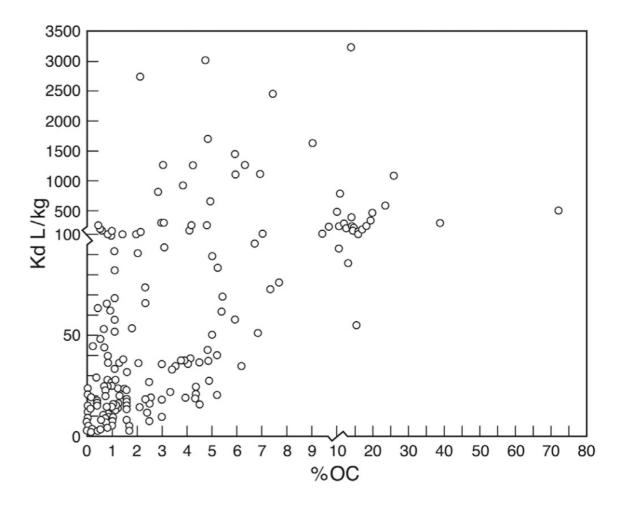


Fig. 1. Relationship between peer-reviewed literature K_d values for PFOS and organic carbon ($R^2 = 0.05$, n = 178).

Source: Li et al, 2018

Field study

To get insight in PFOS leaching at the Oosterweel construction site, a limited field study was performed in which leaching was determined via 2 methods: column leaching and shaking experiments. Unsurprisingly, the field study also revealed a large variability. As is shown on the plot, there is no single curve or line that relates leaching concentration with soil concentration. On the contrary, different lines can be drawn through different subgroups of data. They all correspond to partitioning factors that are in line with Kd values for

PFOS found in international literature, and the variability might reasonably be explained by differences in soil properties at the different sample sites (OC, pH, clay content).

Since there is no data on soil properties covering the total zone of contamination, a safe bet would be to use the leaching data from soils with lowest Kd and use that value to continue our analyses. Note that the data-set is extremely limited, and a safer bet would be to include a safety factor to take into account soils with even lower Kd values that have been missed in this limited field study.

```
# Different K_d values which are all in line with organic matter content betwee 0 and 2%
Kd_1 < -3.5 \# c_b/c_w
Kd_2 <- 12
Kd_3 <- 50
plot(dat$C_s[dat$type=="KP"], dat$C_w[dat$type=="KP"], pch=20, xlab=expression(paste("Bodem [", mu, "g/"
points(dat$C_s[dat$type=="SP"], dat$C_w[dat$type=="SP"], pch=20, col="red")
lines(1:1000, (1:1000)/Kd_1)
lines(1:1000, (1:1000)/Kd_2, lty=2)
lines(1:1000, (1:1000)/Kd_3, lty=3)
legend("top", legend=paste("Kd = ", c(Kd_1, Kd_2, Kd_3), "L/kg"), lty=c(1,2,3), bty="n")
      50
                                             Kd = 3.5 L/kg
                                       ---- Kd = 12 L/kg
      4
                                             Kd = 50 L/kg
Uitloging [µg/L]
      30
      20
      10
                        100
            0
                                     200
                                                 300
                                                              400
                                                                           500
```

State of the art norms for PFOS in soil

Combining our evidence based attribution factor for drinking water, and using the safe bet for the partitioning factor Kd = 3.5, we finally arrive at the state of the art norm or PFOS in soil from the Oosterweel construction site.

Bodem [µg/kg ds]

#

				Soil concen-	Soil concen-	
TDI		drinking water,	drinking water,	tration	tration	
sourceyear (PFOS	S) unit	adult (60kg)	child (10kg)	unit (A)	(C)	$\underline{\text{unit}}$
EFSA2008150	m ng/kg $ m lg/dag$	45	15	ng/L158	52.5	$\frac{-1}{\mathrm{ng/kg}}$
EPA 2009 80	$\frac{ng}{kg}$	24	8	ng/L84	28	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$

TDI sourceyear (PFOS	S) unit	drinking water, adult (60kg)	drinking water, child (10kg)	Soil concentration unit (A)	Soil concentration (C)	unit
EPA 2016 20	ng/kg lg/dag	6	2	ng/L21	7	$\frac{\rm mg/kg}{\rm ds}$
EFSA201820	$\frac{ng}{kg}$	6	2	$\mathrm{ng}/\mathrm{L}21$	7	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$
EFSA2020 0.63	$\frac{ng}{kg}$	0.189	0.063	ng/L0.6	0.2	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$

State of the art norms for PFOS in soil at the time of writing of the RoTS-memo

The RoTS memo was finalized in February 2018. Although there were already European bases exposure pathway studies published in 2011 (e.g. Haug et al., 2011), we can compute what the results would have been when the **best practice WHO attribution factors (0.2)** would have been used

TDI sourceyear (PFOS	S) unit	drinking water, adult (60kg)	drinking water, child (10kg)	Soil concentration unit (A)	Soil concentration (C)	unit
EFSA2008150	ng/kg lg/dag	900	300	ng/L3150	1050	${\mathrm{ng/kg}}$
EPA 2009 80	$\frac{ng}{kg}$	480	160	ng/L1680	560	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$
EPA 2016 20	$\frac{ng}{kg}$	120	40	ng/L420	140	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$
EFSA2018 20	$\frac{ng}{kg}$	120	40	ng/L420	140	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$
EFSA2020 0.63	$\frac{ng}{kg}$	3.78	1.26	ng/L13.23	4.41	$\frac{\mathrm{ng/kg}}{\mathrm{ds}}$

Comparison with RoTS/Lantis value

The difference between the RoTS Lantis value is huge - their derived norm for soil is ... 70.000 ng/kg ds. This is 100.000 to 350.000 times the norm based on best practice and latest evidence and insight, and 165-500 times larger than what would have been best practice at time of writing of the RoTS memo. This is the combined result of cherry picking their "analysis": everywhere possible choices have been made with the most harm to people and planet and the least "obligations" for polluters.

More precisely:

- An outdated value (EFSA, 2008) was used for total daily intake of PFOS. Although the more stringent norms in neighbouring countries and in the US are acknowledged, and the upcoming new EFSA norm (2018) is acknowledged in the memo, nowhere it is motivated why the outdated norm has been chosen. This has resulted in a factor 7 overestimation (at the time of writing) of the TDI. Further, compared to the latest state of the art knowledge EFSA, 2020 norm, a factor 240 overestimation of the TDI was chosen
- Without any motivation, the standard practice of attribution to drinking water (WHO, 2011) was abandoned and 100% of the TDI was allowed through drinking water. At the time of writing of the RoTS memo, this resulted in an additional 5-fold overestimation of allowable concentrations in drinking

water. Taking the current state of the art (where we know that intake via drinking water is $\sim 1\%$ of total intake), the choise for 100% TDI via drinking water results in an additional overestimation of a factor 100.

• From the already limited data set of the field sampling study, the data with largest leaching rates were left out of the analys. This is cherry-picking that does not withstand any good (scientific) practice. The resulting partitioning between pore water and soil underestimates the leaching by an additional factor of about 14.

These 3 cases of cherry picking explain the enormous difference between the RoTS-memo soils concentration of 70.000 ng/kg dw and the evidence based, best practice soil concentration of 0.2-0.6 ng/kg dw.

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