

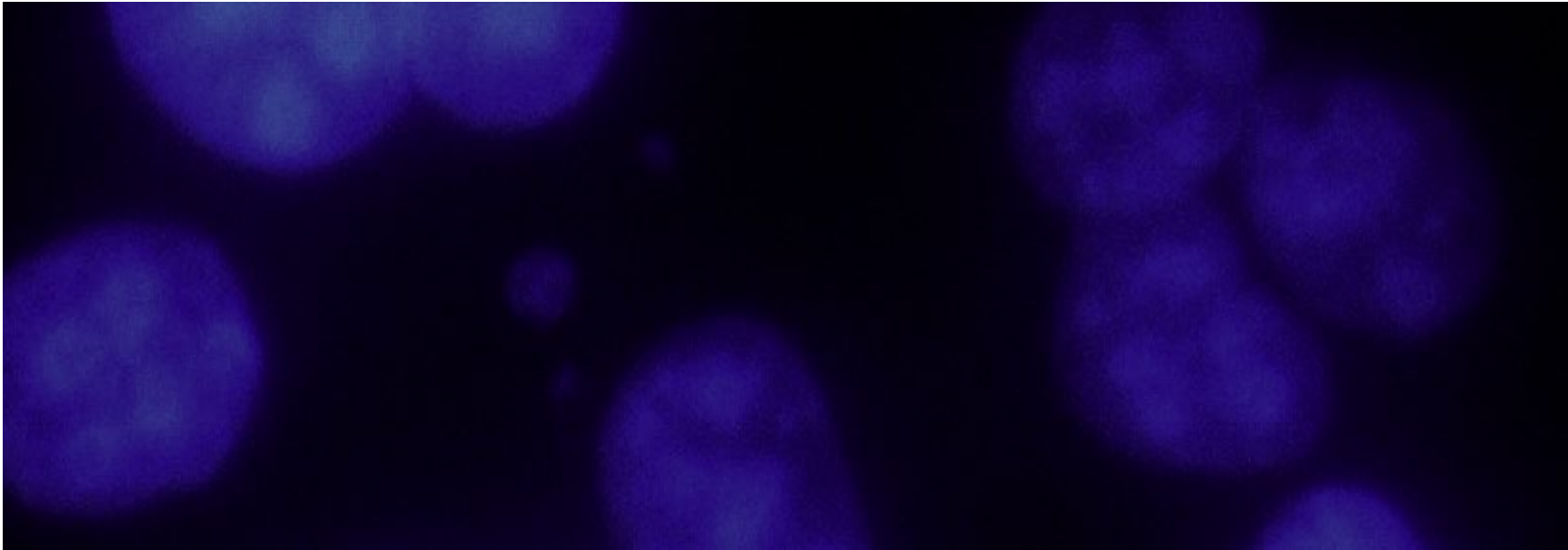
Mixture assessment factors – motivations, derivations and uses

SOT webinar 11 Jan 2023

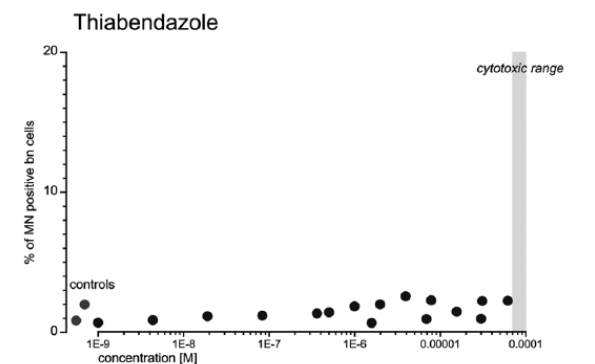
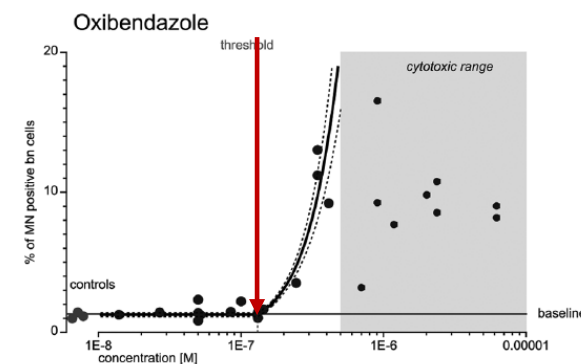
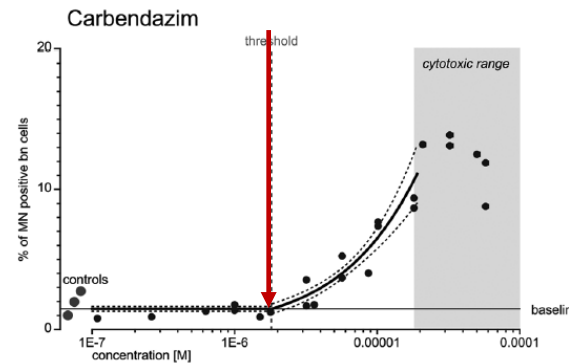
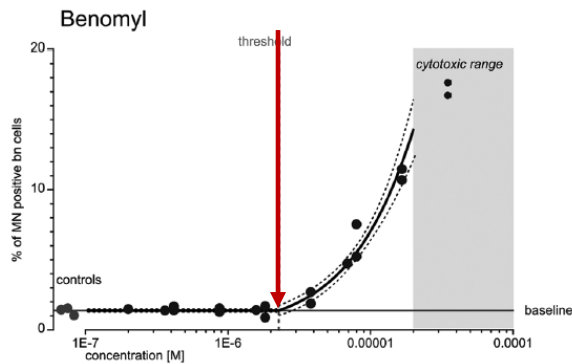
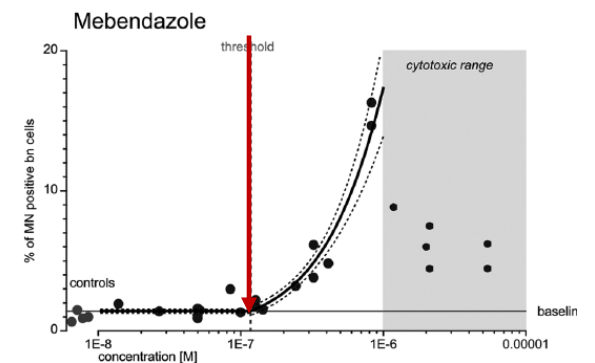
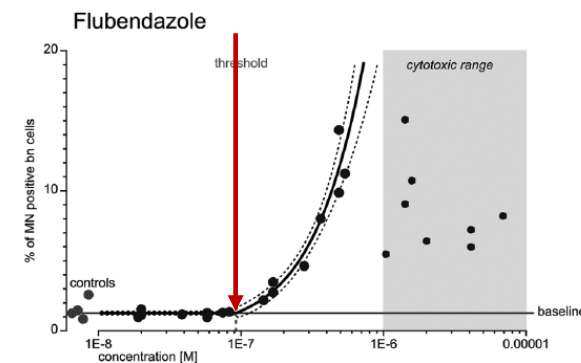
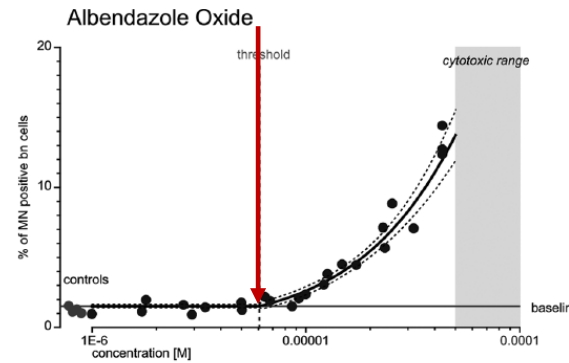
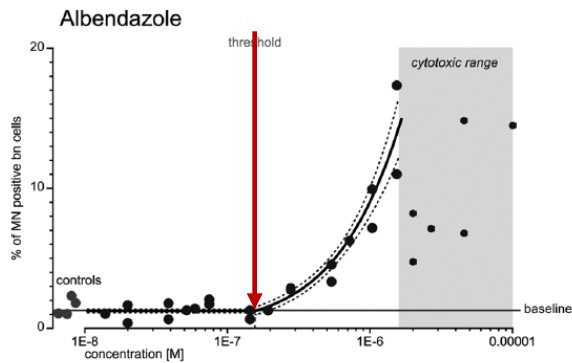
Andreas Kortenkamp, Brunel University London

“Protection from mixture effects is achieved, as long as exposures stay below thresholds”

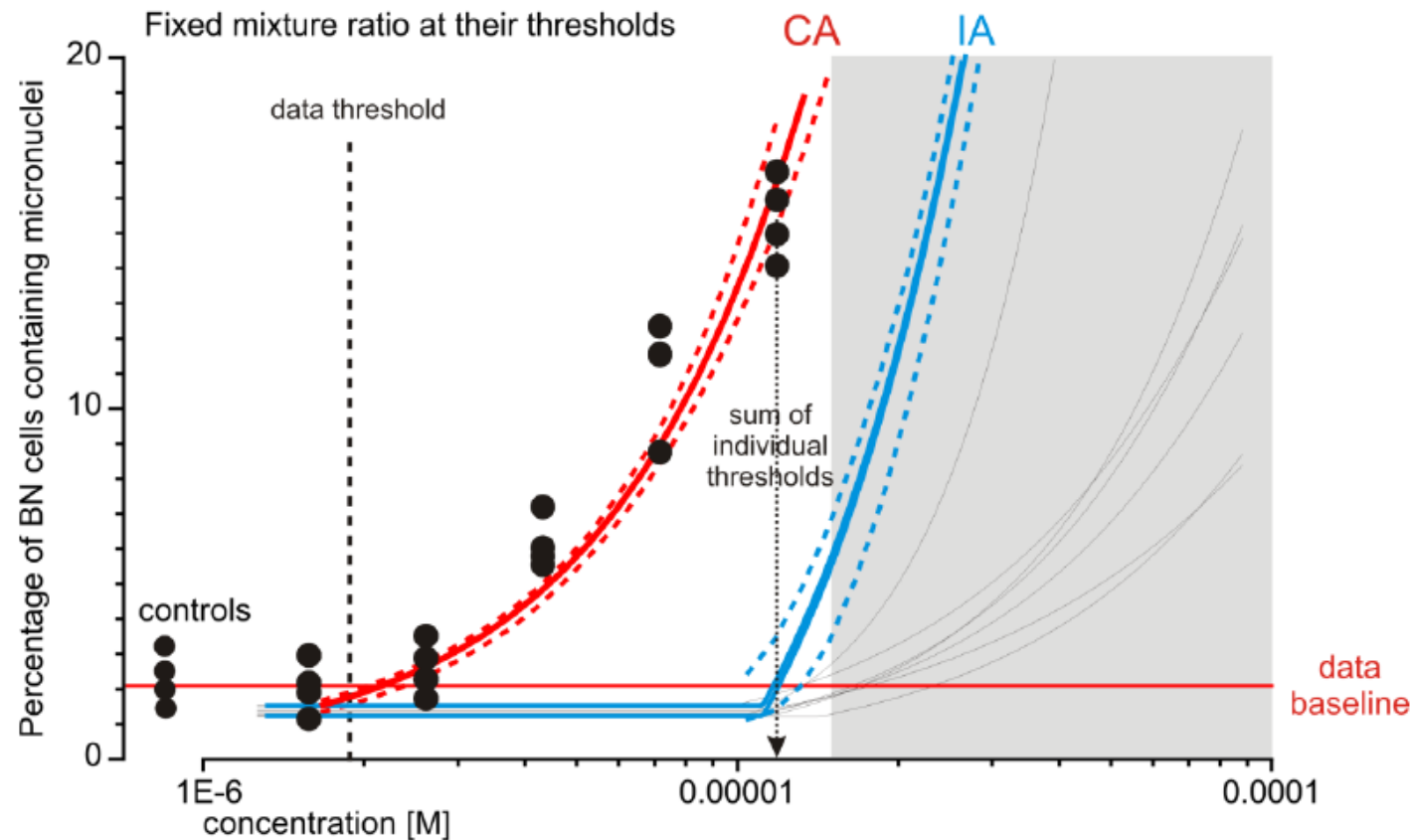
Micronuclei



Benzimidazoles and micronuclei – concentration-response relationships



Benzimidazole low dose mixture: micronuclei



Clear combination effects at
sub-threshold doses

= Dose addition expectation

Seven benzimidazole pesticides combined at sub-threshold levels induce micronuclei *in vitro*

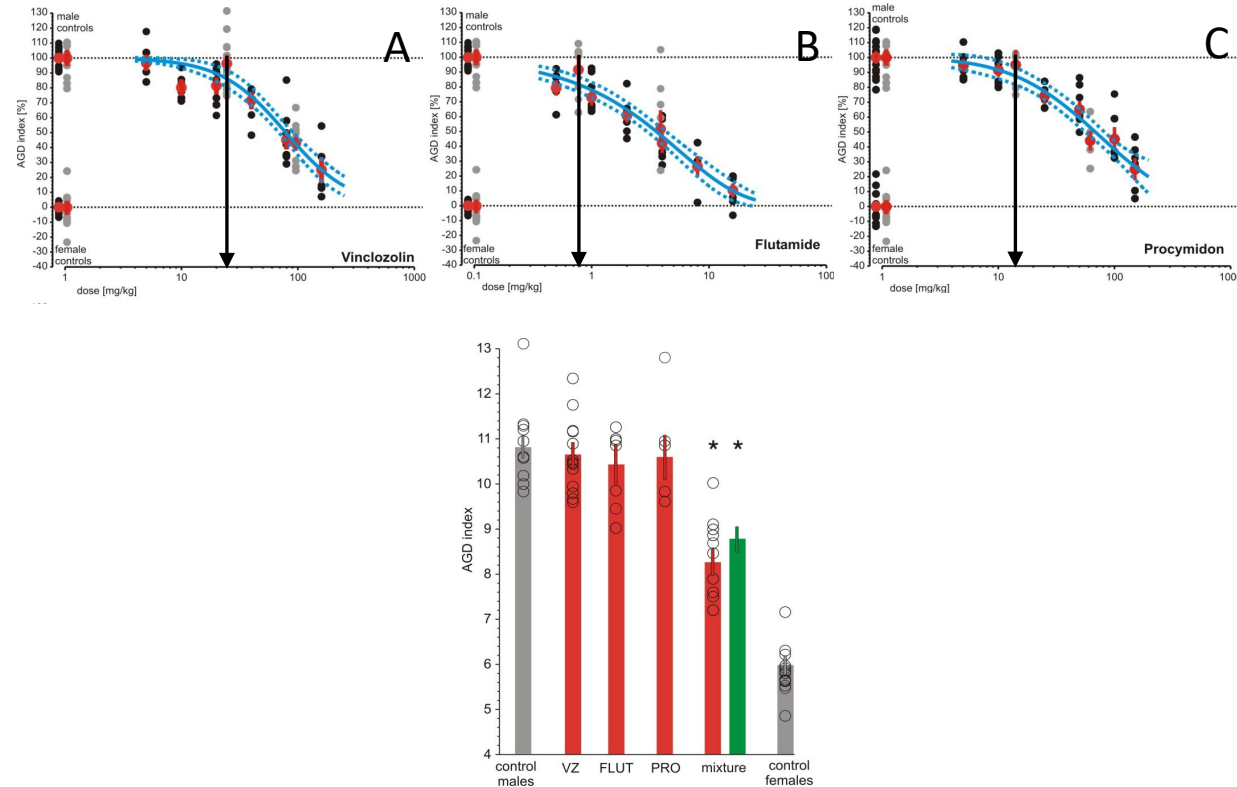
Sibylle Ermler*, Martin Scholze and Andreas Kortenkamp

Institute for the Environment, Brunel University, Kingston Lane, Uxbridge,
Middlesex UB8 3PH, UK

*To whom correspondence should be addressed. Tel: +44 1895 267208;
Fax: +44 1895 268761; Email: sibylle.ermler@brunel.ac.uk

colchicine-binding site (1). In target organisms, the intended effect is cytotoxicity, which occurs through disruption of microtubuli (2,3). At lower, non-cytotoxic concentrations, the impairment of the microtubuli of the spindle apparatus can disturb the alignment of chromosomes during mitosis and lead to the formation of micronuclei (MN). In cultured mammalian

Anti-androgens in a rat developmental toxicity model



Hass U, Scholze M, Christiansen S, Dalgaard M, Vinggaard AM, Axelstad M, Metzdorff SB, Kortenkamp A: **Combined exposure to anti-androgens exacerbates disruption of sexual differentiation in the rat.** *Environ Health Perspect* 2007, 115(Suppl 1):122-128.

Mixture effects at low doses,
below effect thresholds

Possible implications

Compliance with single chemical regulatory exposure limit values **not necessarily protective**

Lower limit values for protection against mixture risks?

Mixture risk assessment case study

- Scale of the problem
- Mitigation
- **Mixture assessment factors** as a solution?

Application to risk assessment practice

Hazard Index

Sum of “risk quotients”

$$\frac{\text{Intake}_1}{\text{Reference dose}_1} + \frac{\text{Intake}_2}{\text{Reference dose}_2} < 1$$

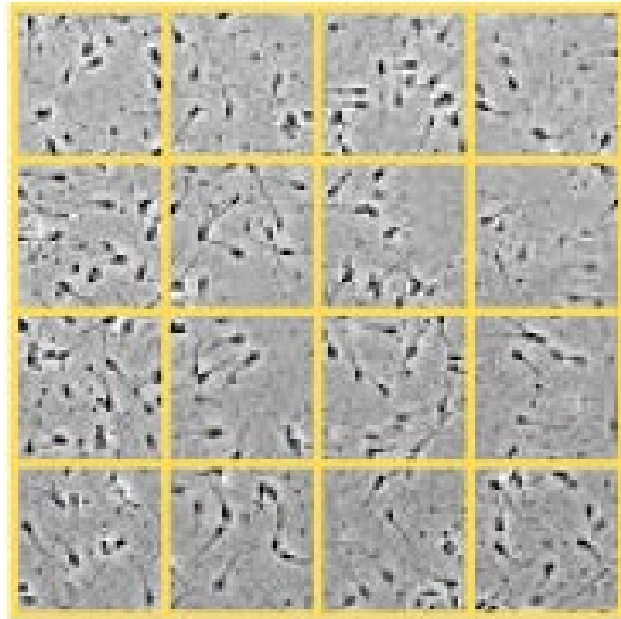
- **Assumptions**

- Dose addition as a good approximation
- No synergisms

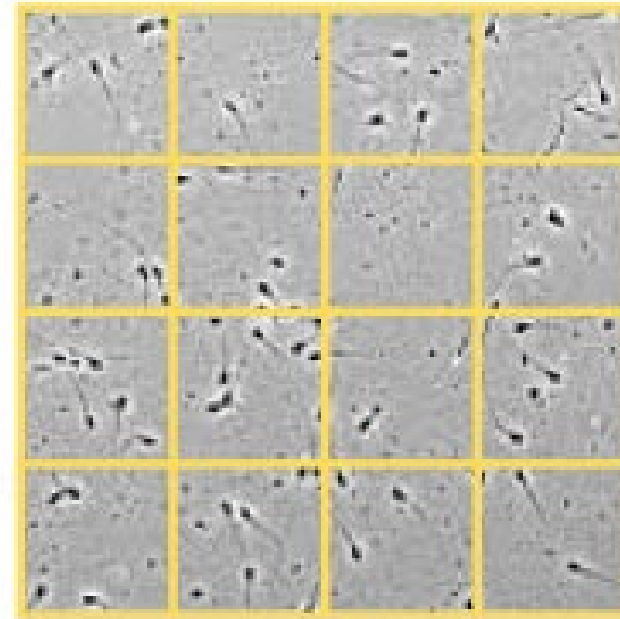
- **Interpretation**

- Exceedance of combined acceptable **exposures**
- NOT estimation of effect size!

Sperm concentration



101 mill/ml
(1973)



49 mill/ml
(2018)

29 chemicals

PCDD/F			Phthalates	DiBP
PBDE	BDE-209			DnBP
	BDE-183			DEHP
	BDE-154			DiNP
	BDE-153			BBzP
	BDE-100		Bisphenols	BPA
	BDE-99			BPS
	BDE-47			BPF
	BDE-28		Painkillers	Paracetamol
PCB	PCB 169			
	PCB 126			
	PCB 118			
Acrylamide				
n-Butyl paraben				
Pesticides	Chlorpyrifos			
	Vinclozolin			
	Procymidone			
	Prochloraz			
	Linuron			
	Fenitrothione			

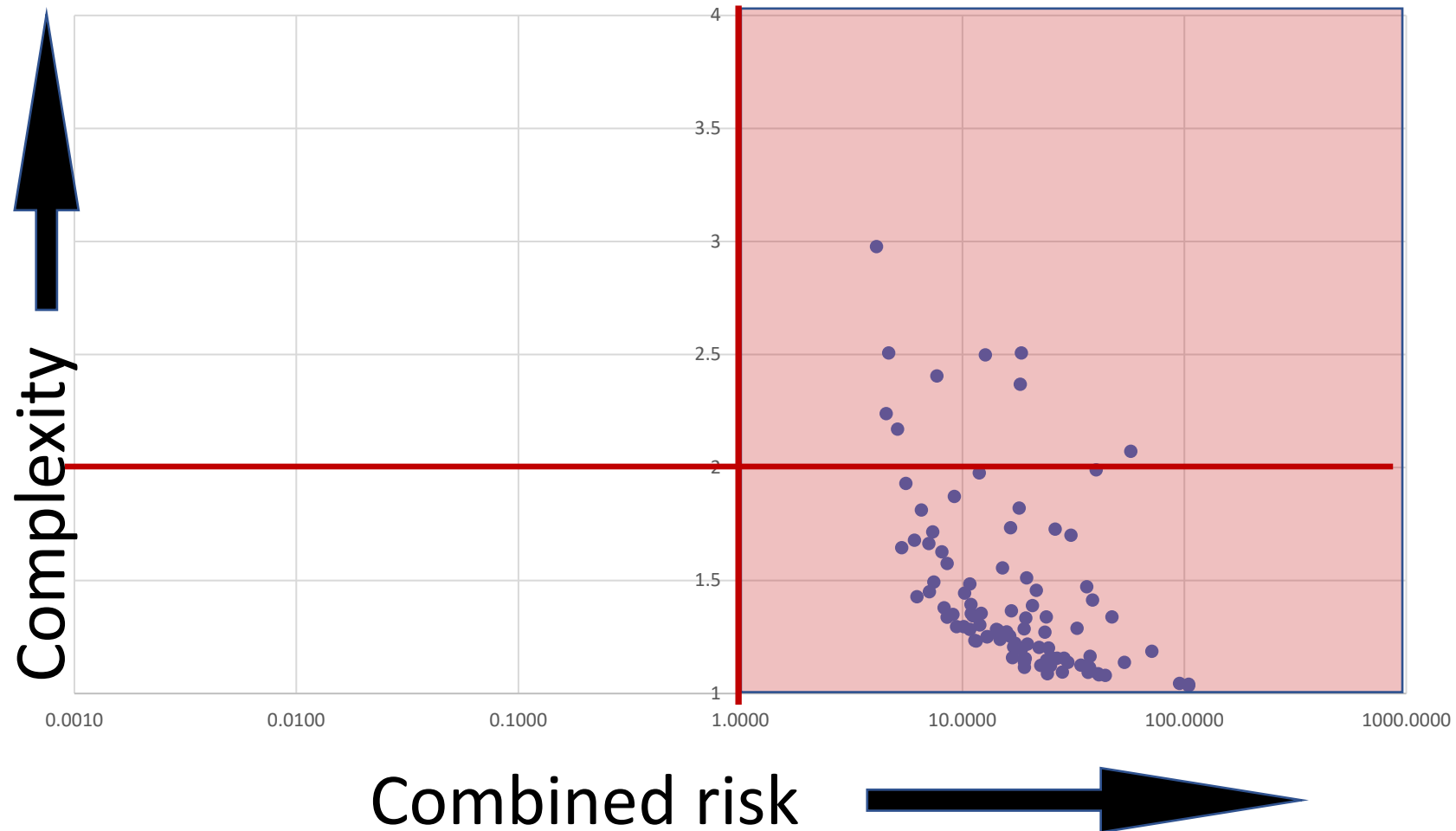
Multiple chemicals monitored in the same sample



Phthalates	DiBP
	DnBP
	DEHP
	DiNP
	BBzP
Bisphenols	BPA
	BPS
	BPF
Painkillers	Paracetamol

Mixture risk assessment gets personal

Phthalates, bisphenols, paracetamol + 20 background chemicals¹⁰

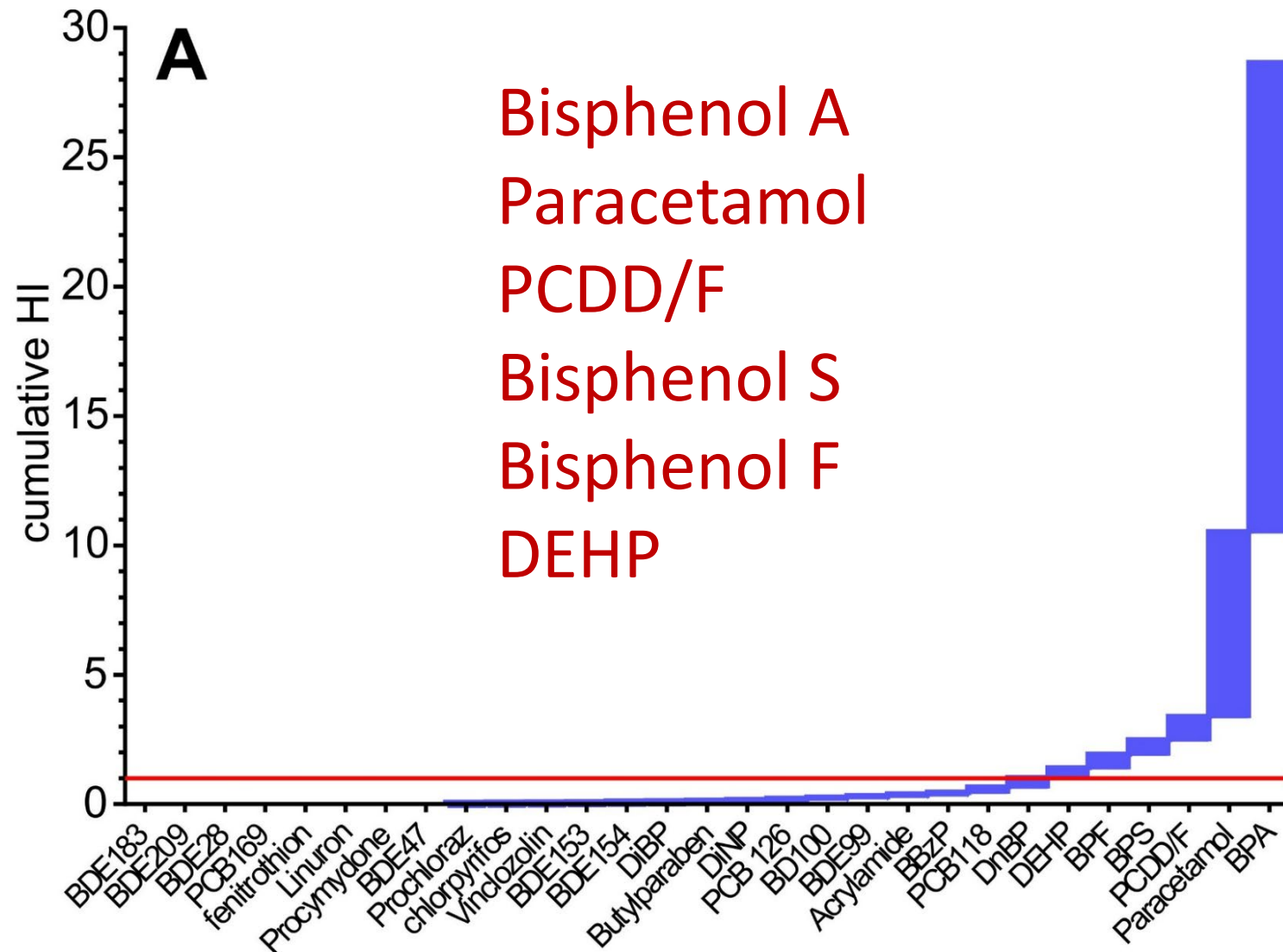


Exceedances of combined “acceptable” exposures

Range: 4 – 100-fold

Median: 18-fold

Drivers of mixture risks (semen quality)



Exceedance if all risk quotients ≤ 1

\sim 5-fold



Contents lists available at [ScienceDirect](#)

Environment International

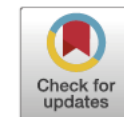
journal homepage: www.elsevier.com/locate/envint



Full length article

Combined exposures to bisphenols, polychlorinated dioxins, paracetamol, and phthalates as drivers of deteriorating semen quality

Andreas Kortenkamp^{a,*}, Martin Scholze^a, Sibylle Ermler^a, Lærke Priskorn^{b,c},
Niels Jørgensen^{b,c}, Anna-Maria Andersson^{b,c}, Hanne Frederiksen^{b,c}



Exposure limits for single chemicals **do not protect** against mixture risks

Assessment factors used to derive limit values do not deal with mixture risks

An additional factor is needed: **Mixture Assessment Factor (MAF)**

Two uses of MAFs

1. For downward correction of exposure limits

(regulatory values, reference doses, ADI, TDI)

Mixture **Assessment** Factor



2. For downward correction of index values in risk assessment

Risk quotient = 1

Risk quotient = 0.1; 0.01 etc

Mixture **Allocation** Factor



Proposed MAFs (Europe)

Area	Size	Reference
Environment	100	Janssen, 2004; van Vlaardingen, 2007
Environment	100	Tørsløv, 2013
Human health	100	Muilerman, 2011
Human health	10	Tørsløv, 2013, Petersen, 2014

No justifications given

Mode of application not defined

MAFs: an alluring solution?

- Easy to understand
- Easy to use
- A pragmatic approach to a complicated problem

MAFs: the criticism

- Arbitrary
- Not science-based
- Not data-driven

Theory- and data-driven sizing of a MAF

When are mixed exposures “safe”?

$$HI = \sum_{i=1}^n \frac{EL_i}{AL_i} \leq 1$$

HI = Hazard Index; EL = Exposure Level; AL = Acceptable Level (RfD)

No exceedance of HI = 1 if:

$$EL = 1/n \times AL$$

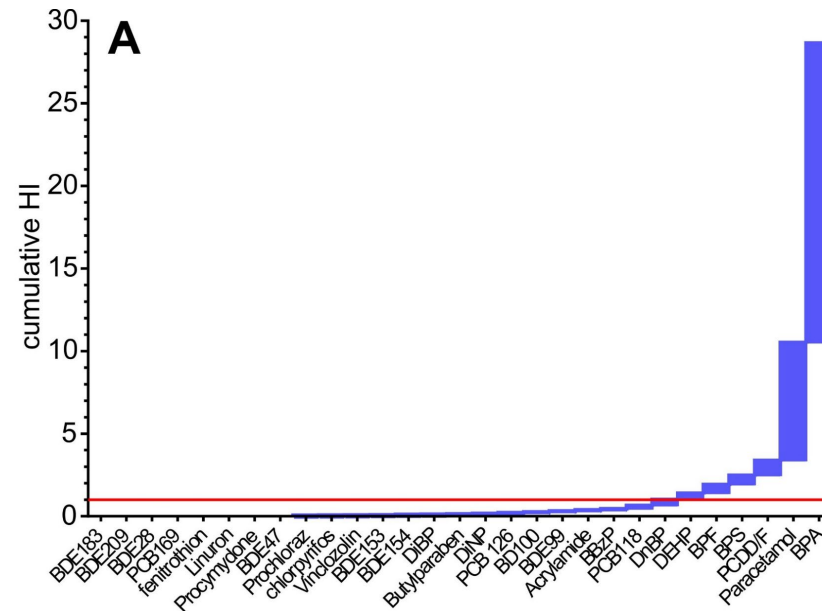
n = number of mixture components

Lower each AL by a factor of **1/n**

n = Mixture **Assessment** Factor

MAF = n overly conservative

Smaller values also meet condition $HI \leq 1$ when risk quotients are unevenly distributed



Data-driven sizing of a Mixture **Assessment** Factor



Mixture **Assessment** Factor

$$\sum_{i=1}^n \frac{EL_i}{AL_i} \times \frac{1}{MAF} \leq 1$$

KEMI (2015)

Procedure:

1. To separate single chemical compliance issues from true mixture problems, **set all RQ > 1 to 1**
2. Calculate **sum of adjusted RQ**
3. Use sum of adjusted RQ as **Mixture **Assessment** Factor**

Mixture **Assessment** Factor: Example

Lower bound, based on P50 or geometric mean, 2009 exposures					
Chemical	Exposure	Unit	Reference dose	Unit	Risk Quotient
Bisphenol A	0.048	µg/kg d	0.01	µg/kg d	1
Paracetamol	7	mg/kg d	1	mg/kg d	1
PCDD/F	0.25	pg/kg d	0.28	pg/kg d	0.9
Bisphenol F	0.006	µg/kg d	0.01	µg/kg d	0.6
DEHP	2.06	µg/kg d	10	µg/kg d	0.206
Bisphenol S	0.002	µg/kg d	0.01	µg/kg d	0.2
PCB 118	575	pg/kg/d	2900	pg/kg/d	0.198
DBP	0.88	µg/kg d	6.7	µg/kg d	0.131
BDE 99	0.18	ng/kg/d	2.88	ng/kg/d	0.063
BDE 100	0.15	ng/kg/d	2.88	ng/kg/d	0.052
Acrylamide	0.4	µg/kg d	8.3	µg/kg d	0.048
PCB 126	3.5	pg/kg/d	73	pg/kg/d	0.048
n-butylparaben	0.6	µg/kg d	30	µg/kg d	0.02
BDE 154	0.05	ng/kg/d	2.88	ng/kg/d	0.017
BBzP	0.15	µg/kg d	10	µg/kg d	0.015
BDE 153	0.04	ng/kg/d	2.88	ng/kg/d	0.014
DINP	0.77	µg/kg d	59	µg/kg d	0.013
DIBP	0.99	µg/kg d	100	µg/kg d	0.010
Vinclozolin	0.35	µg/kg d	50	µg/kg d	0.007
BDE 47	0.58	ng/kg/d	150	ng/kg/d	0.004
Procymidone	0.25	µg/kg d	100	µg/kg d	0.0025
Prochloraz	0.34	µg/kg d	160	µg/kg d	0.002
Linuron	0.069	µg/kg d	100	µg/kg d	0.00069
PCB 169	3.5	pg/kg/d	5330	pg/kg/d	0.00066
Fenitrothione	0.06	µg/kg d	200	µg/kg d	0.0003
BDE 28	0.02	ng/kg/d	150	ng/kg/d	0.000
BDE 209	0.61	ng/kg/d	1000000	ng/kg/d	0.000
BDE 183	0.02	ng/kg/d	1000000	ng/kg/d	0.000
Sum of RQ					4.55
MCR					4.55

n = 29

Mixture **Assessment** Factor: 4.55

Compliance with **4.55-fold lower Reference Doses** achieves $HI \leq 1$

Lowering by a factor of 29 is not necessary



**So
unfair!**

Data-driven sizing of a Mixture **Allocation** Factor



Mixture **Allocation** Factor

$$\sum_{i=1}^n \frac{EL_i}{AL_i} > \frac{1}{MAF} = \frac{1}{MAF} \leq 1$$
$$\sum_{i=1}^n \frac{EL_i}{AL_i} \leq \frac{1}{MAF} = \frac{EL_i}{AL_i} \leq 1$$

KEMI (2021)

Procedure:

1. To separate single chemical compliance issues from true mixture problems, **set all RQ > 1 to 1**
2. Through **iteration**, determine 1/MAF so that **sum of adjusted RQ = 1**

Mixture **Allocation** Factor: Example

	Risk Quotient	Risk Quotient adjusted	Risk Quotient with MAF										
DiBP	0.007	0.007	0.007										
DnBP	0.231	0.231	0.139	If Risk Quotient adjusted $\leq 1/\text{MAF}$: no change									
BBzP	0.017	0.017	0.017										
DEHP	0.209	0.209	0.139	If Risk Quotient adjusted $> 1/\text{MAF}$: change to $1/\text{MAF}$									
DiNP	0.006	0.006	0.006										
BPF	3.140	1.000	0.139	Aim: Sum of Risk Quotients = 1									
BPS	0.332	0.332	0.139										
BPA	7.363	1.000	0.139										
Paracetamol	5.774	1.000	0.139										
PCDD/F	0.893	0.893	0.139										
		4.7	1.0										
Mixture Allocation Factor	7.2												
inverse	0.139												

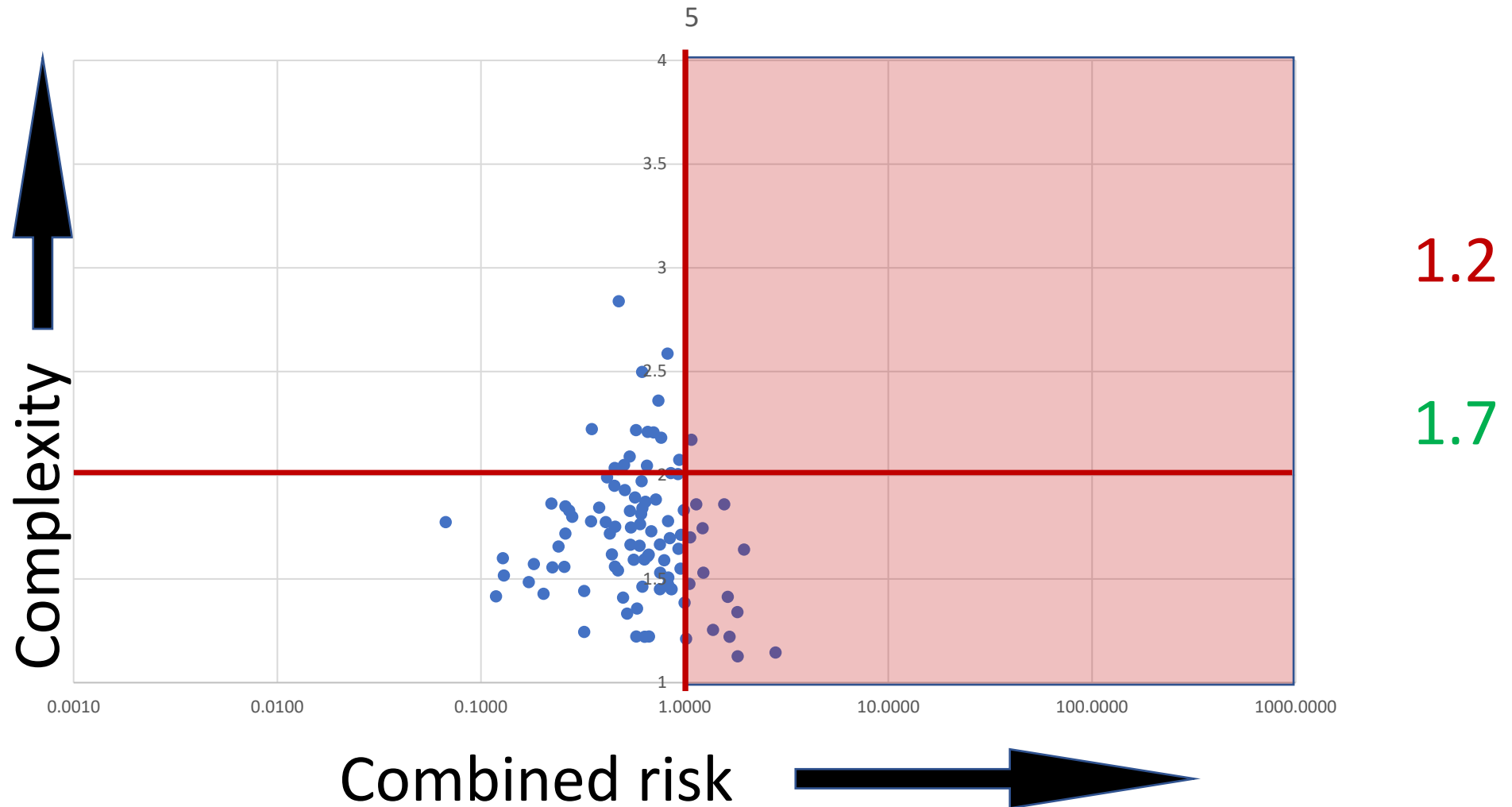
Small Risk Quotients are left **untouched**

The price: MAF has to be **larger** (here: 7.2 versus 4.7)

A scientifically sound data-driven approach to sizing a MAF requires **comprehensive information** about relevant exposure scenarios

Five phthalates

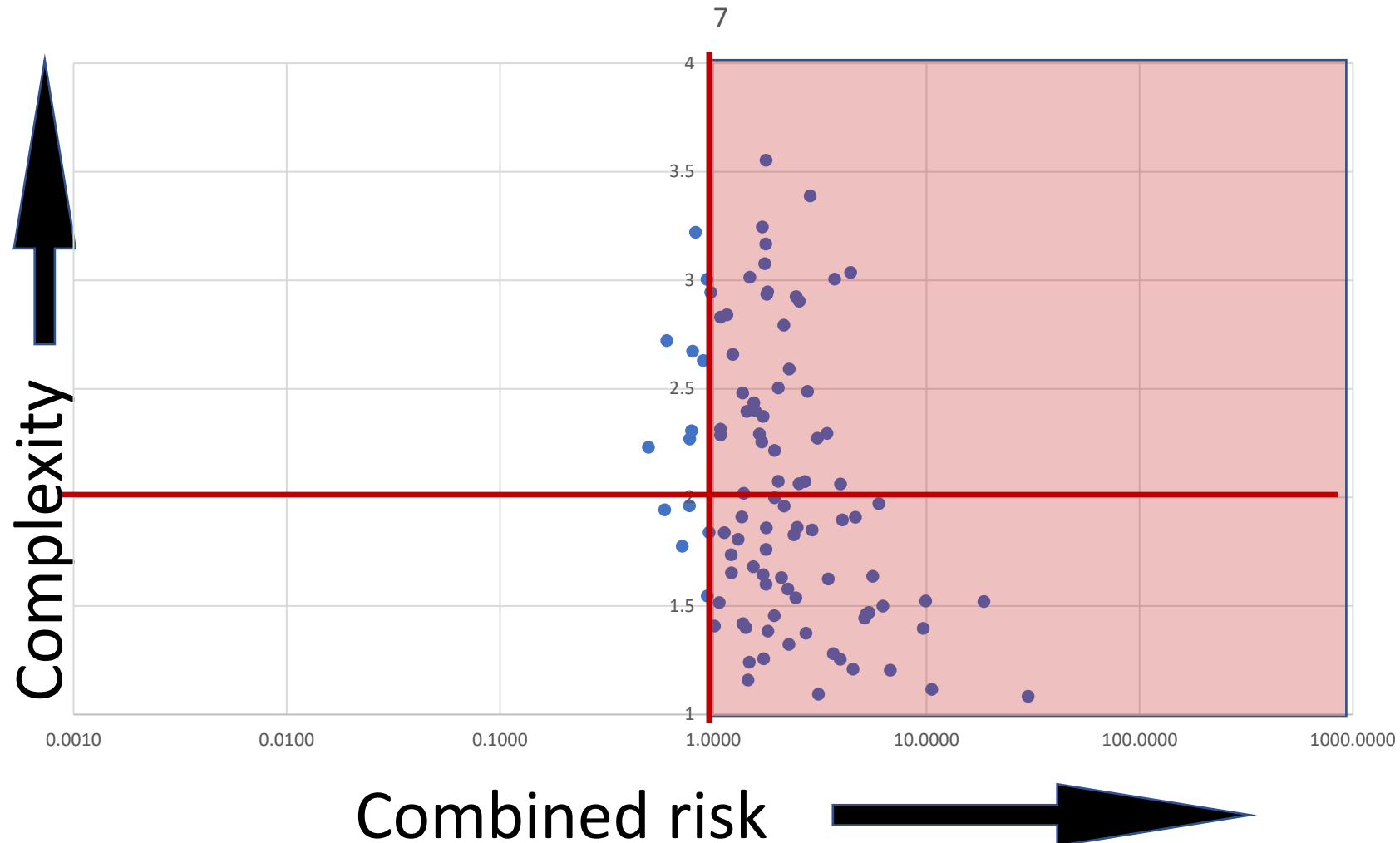
Mixture **Assessment** factor: 1.2
Mixture **Allocation** Factor: 1.7



Five phthalates, bisphenol F, S

Mixture **Assessment** factor: 2.2

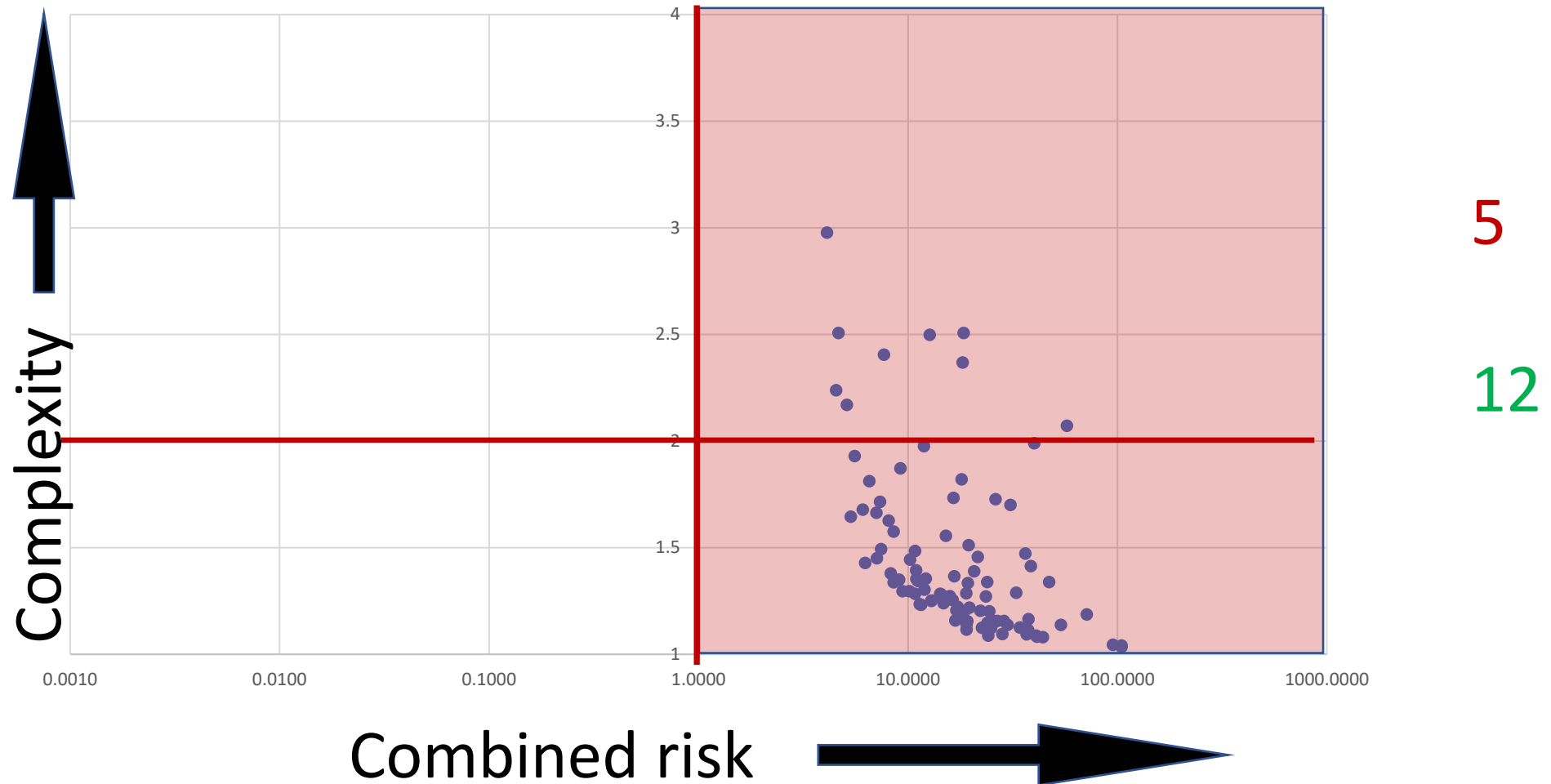
Mixture **Allocation** Factor: 4



Phthalates, bisphenols, paracetamol + 20 background chemicals

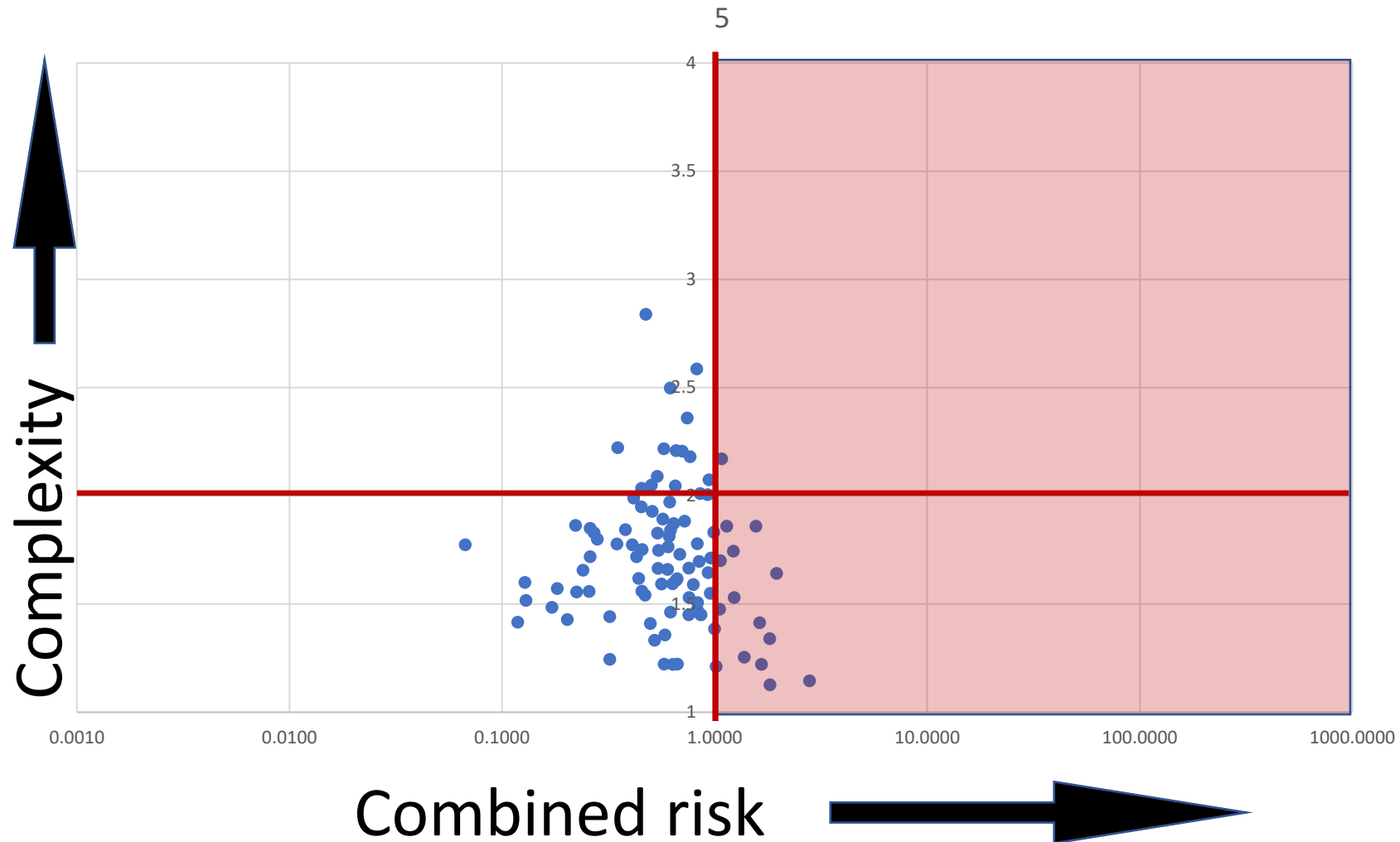
Mixture **Assessment** factor: 5

Mixture **Allocation** Factor: 12

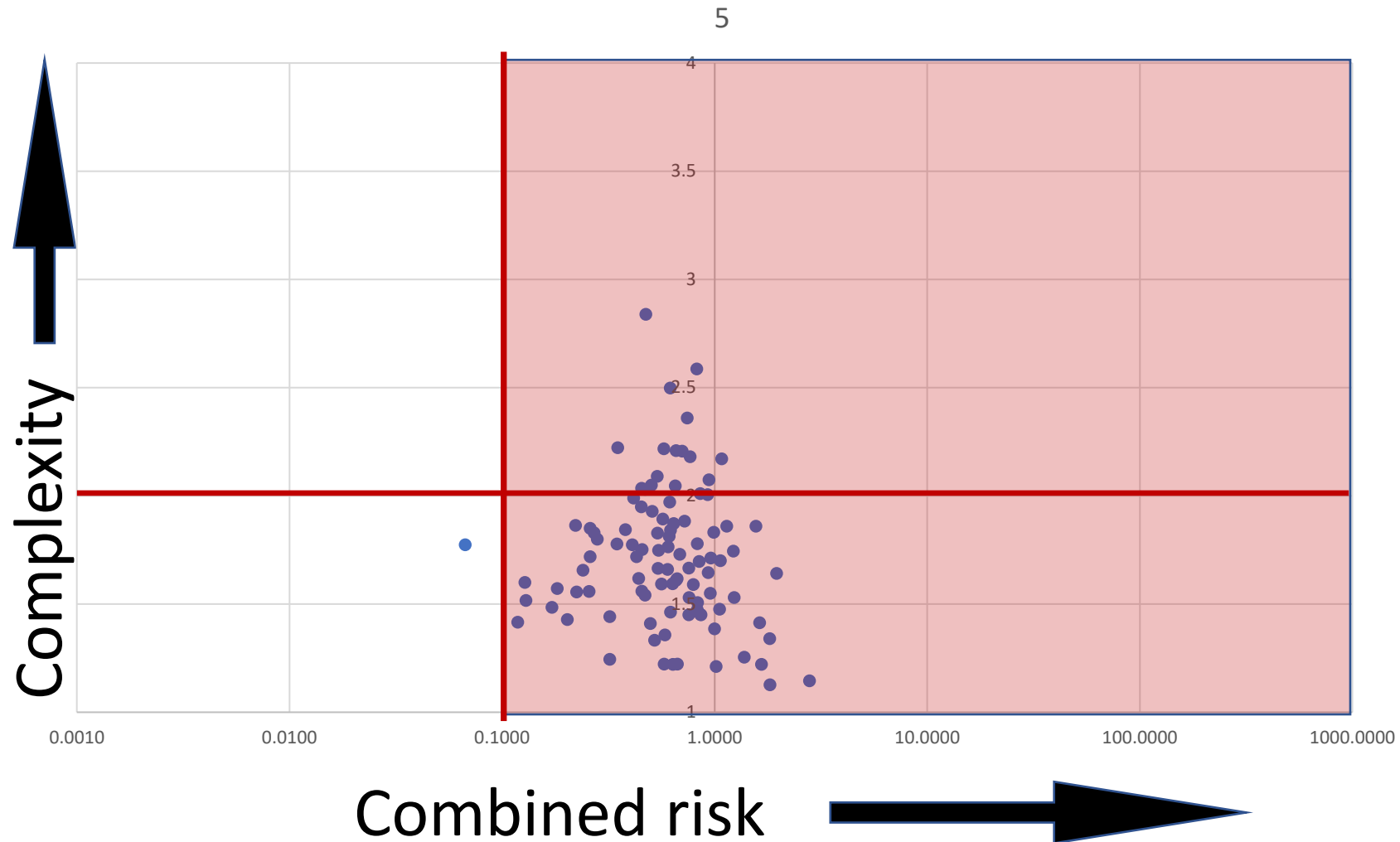


With incomplete exposure information,
but a **more cautious approach** to risk
assessment –
would we make better decisions?

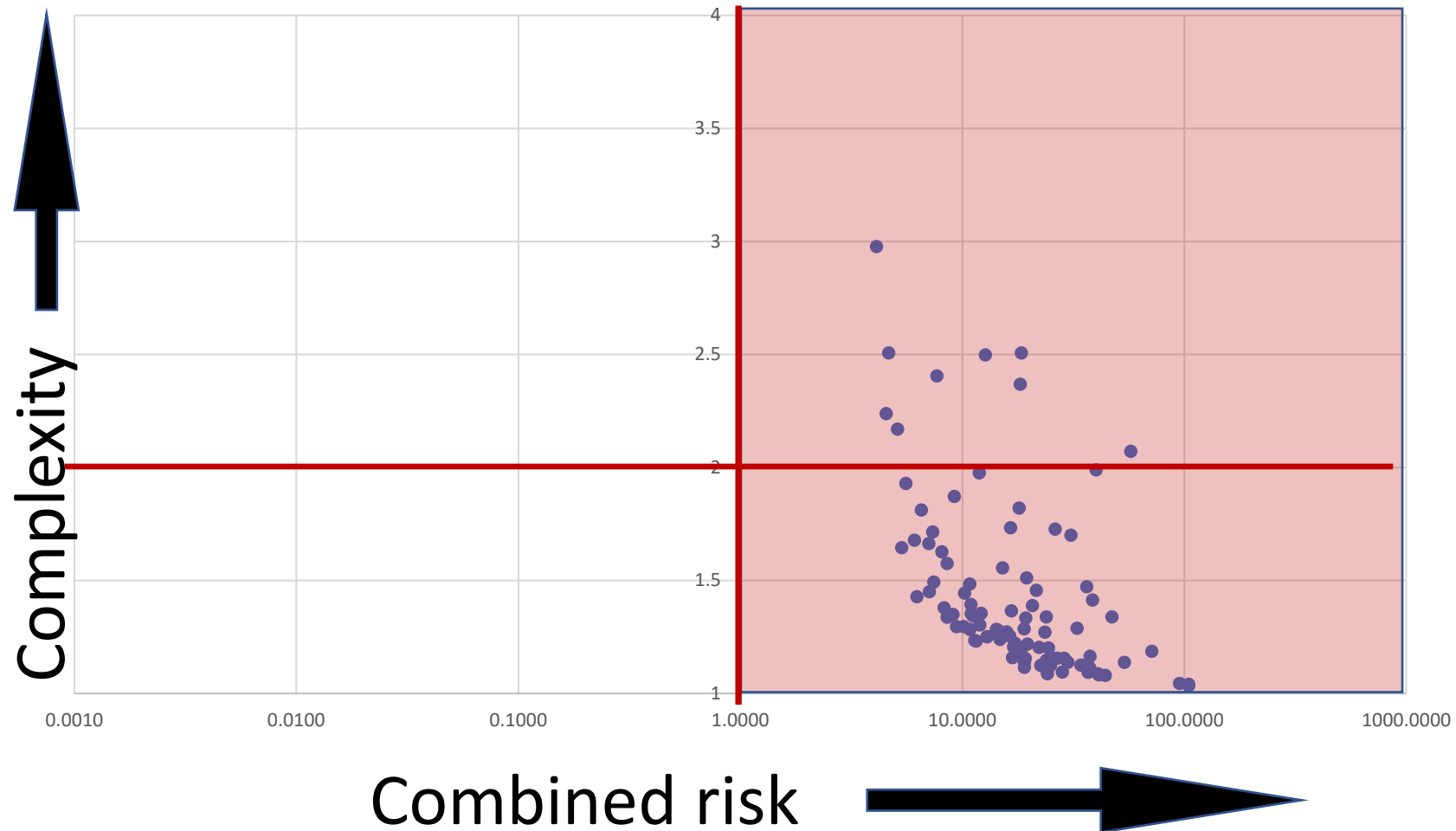
Five phthalates: **MAF = 1**



Five phthalates: **MAF = 10**



Phthalates, bisphenols, paracetamol + 20 background chemicals¹⁰



This case study supports:

Mixture **Assessment** factor: **5**

Mixture **Allocation** Factor: **12**

BUT:

MAF > 5 not achievable in the context of
REACH (industrial chemicals)

Residual risk with Mixture

Allocation Factor = 5

2-fold exceedance of acceptable
combined exposures

Residual risk with Mixture

Assessment Factor = 5

No exceedance of acceptable
combined exposures

However:

Minimum risk estimate

Not taken into account:

- Air pollution
- Perfluorinated chemicals

Risk estimates and **MAFs increase** the more substances are included in the assessment

Conclusions

- Use of a MAF in risk assessment and risk management is **scientifically justified**
- **Practicable**: Can be integrated in current risk management approaches
- **Urgent** for the protection against mixture risks
- More human health case studies needed to support data-driven sizing of a MAF

**Thank
you!**

