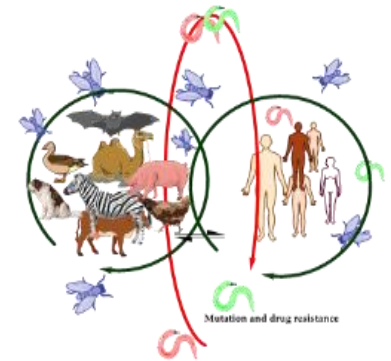


Environmental impact of Pharmaceuticals

Rolf-Alexander Düring
Institute of Soil Science and Soil Conservation
Justus Liebig University Gießen



One Health drugs against
parasitic vector borne diseases in
Europe and beyond -
OneHealthdrugs

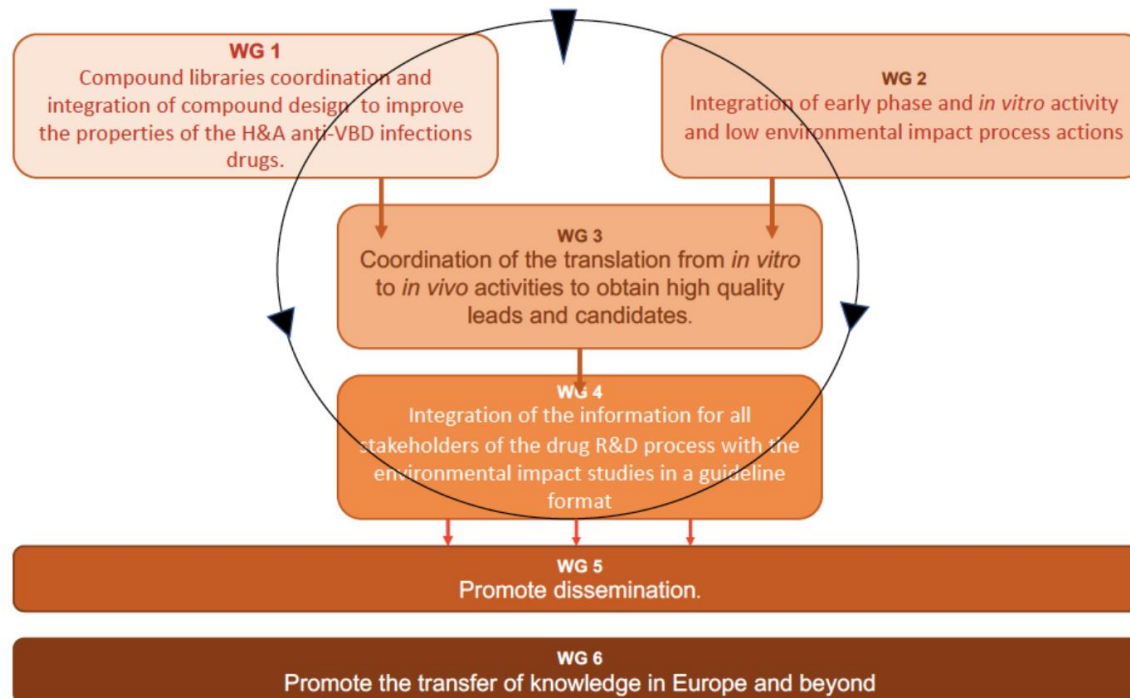


WG4, Workshop 1:
Environmental impact of
pharmaceuticals and
international organizations
monitoring
03/04/2023



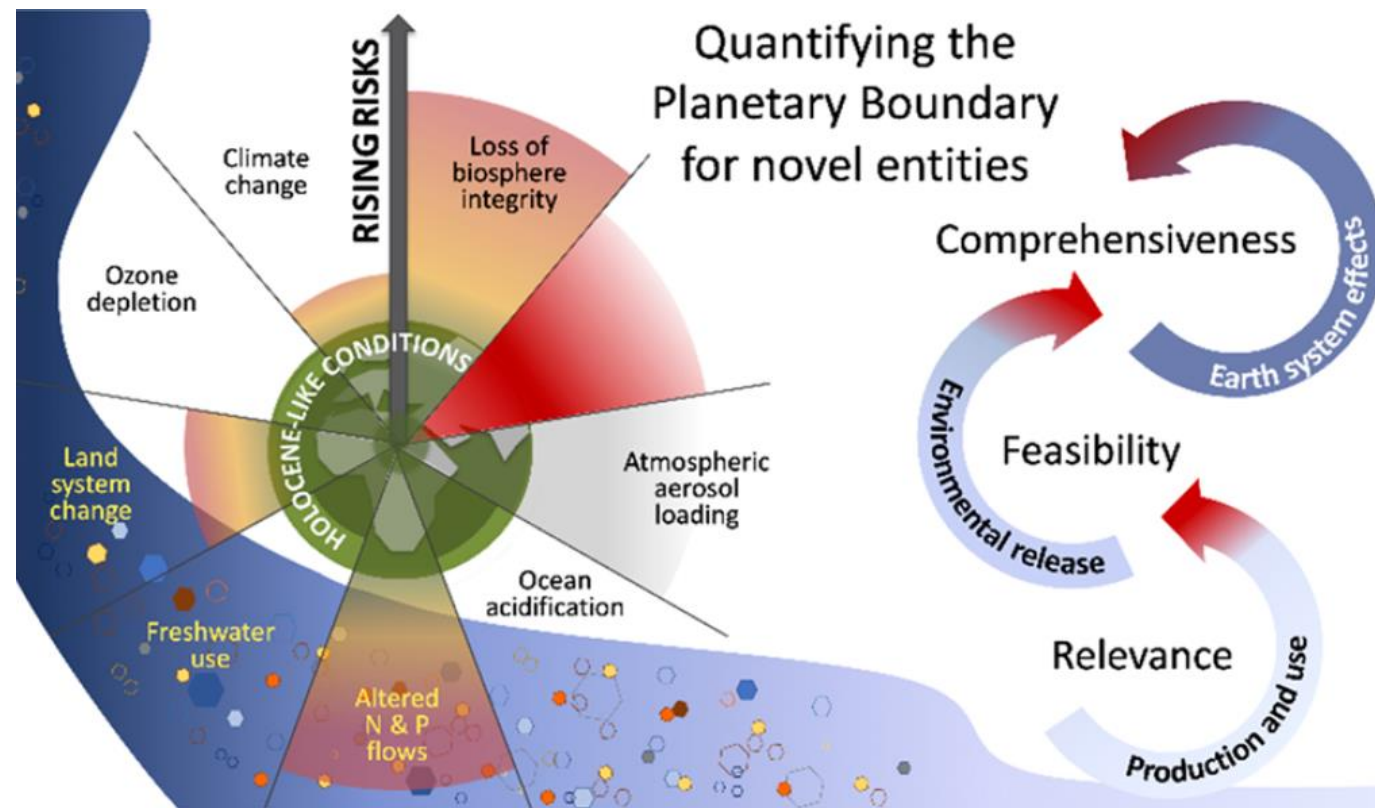
WG4: Integration of R&D Process-Environmental Studies for the Translation in a White Paper

- Drug design in compliance with the overall environmental impact to provide a sharable guideline-like document.
- Assessment of probability of exposure on the basis of substances environmental fate.
- Properly inform drug designers and managers on environmental risks compared to societal benefits.



Are we leaving the Safe Area of the Planetary Boundary for Novel Entities?

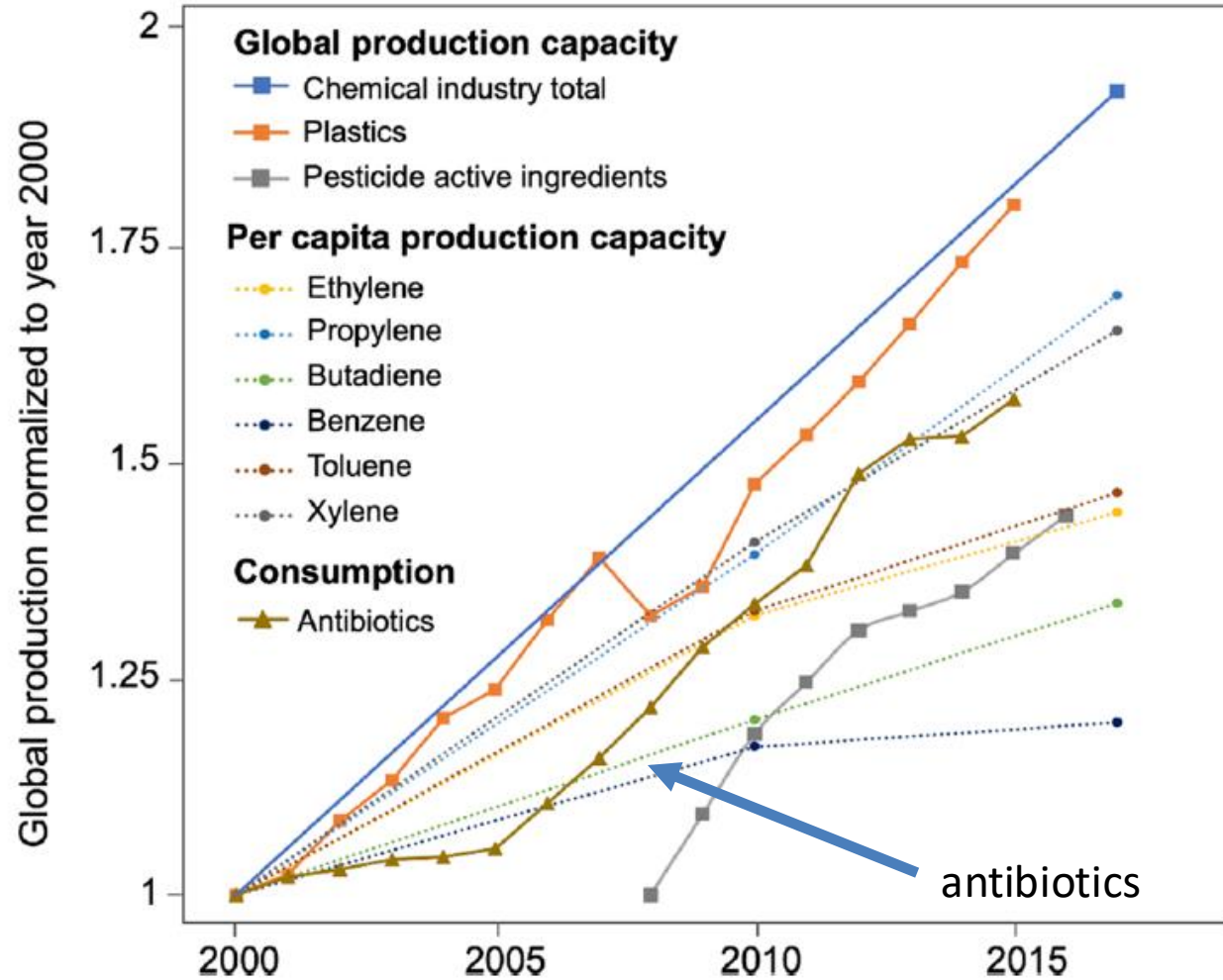
“...the anthropogenic introduction of novel entities to the environment is of concern at the global level when these entities exhibit persistence, mobility across scales with consequent widespread distribution and accumulation in organisms and the environment, and potential negative impacts on vital Earth System processes or subsystems...”



(Persson et al., 2022)

Relative Growth in the Production of Certain Chemicals

- Polymer "plastic" as the most visible form of chemical pollution: total mass of plastic exceeds that of all living mammals
- 2.4% of it enters the environment every year

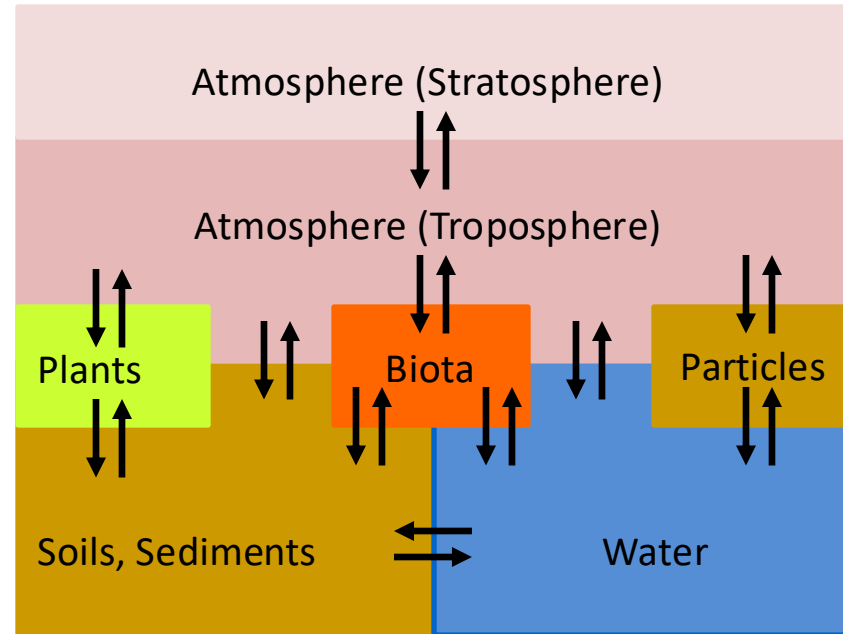


(Persson et al., 2022)

Environmental Impact: What is it all about?

what does the environment do to the substance?

Environmental Chemistry:
Partitioning, transformation, exposure



what does the substance do to the environment?

(Eco)toxicology:
Effects on non-target organisms

nature International weekly journal of science

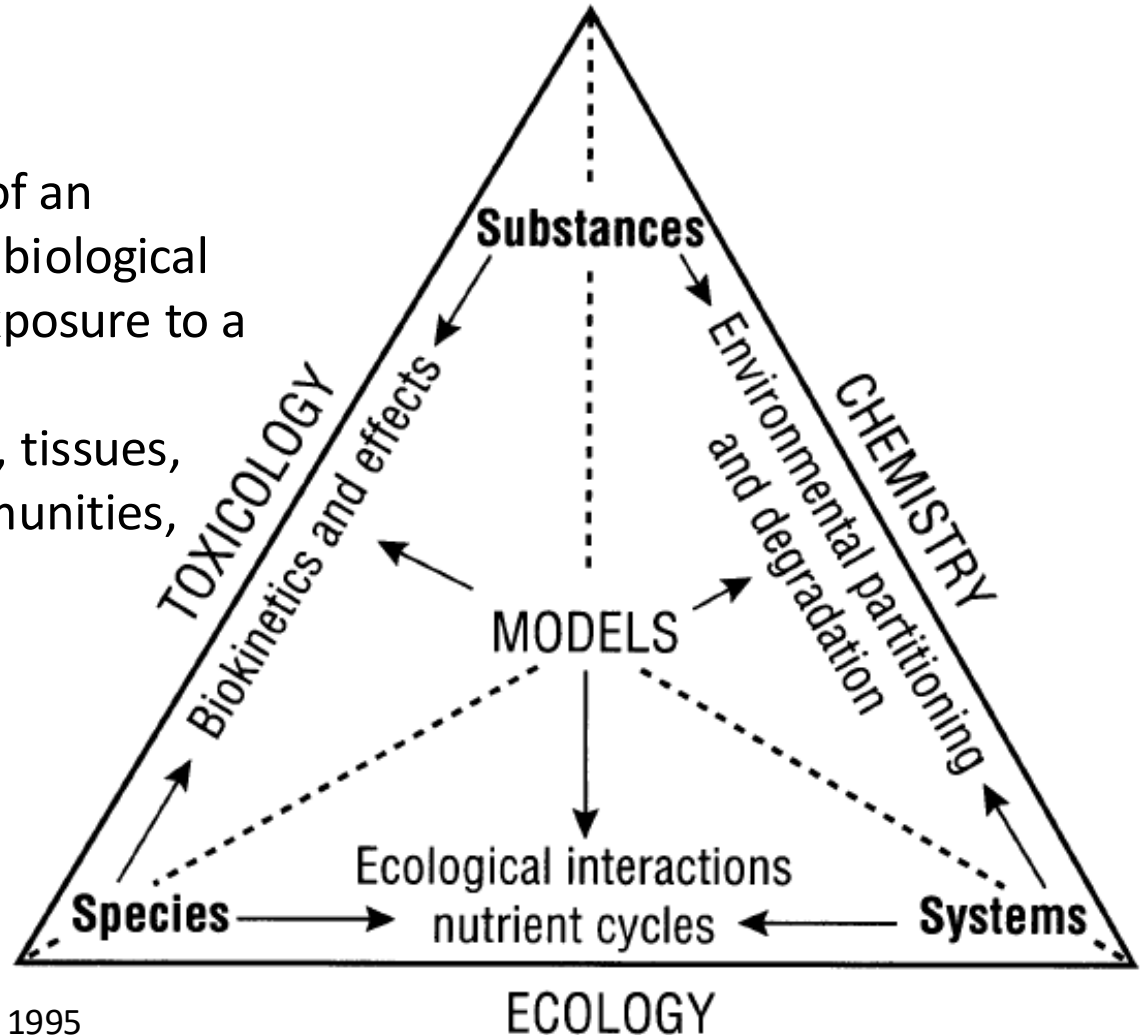
**Diclofenac residues as the
cause of vulture population
decline in Pakistan**

Oaks et al. (2004)

Ecotoxicology: “Study of the three S's”

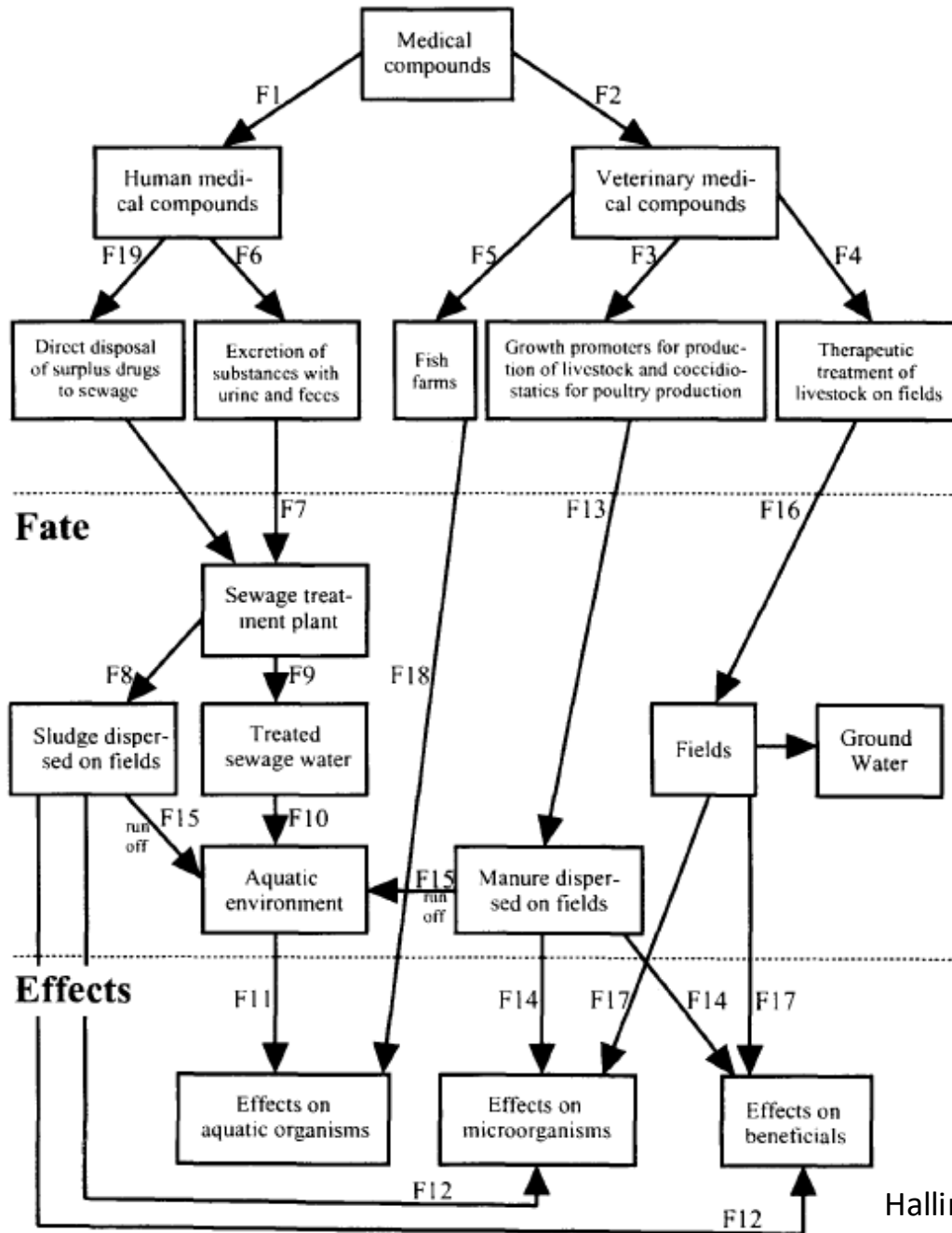
Toxic Effects of **S**ubstances on Nonhuman **S**pecies in Complex **S**ystems

- Changes in state or dynamics of an organism, or at other levels of biological organization, resulting from exposure to a chemical
- From subcellular/cellular level, tissues, individuals, populations, communities, ecosystems, and landscapes



Van Leeuwen, 1995

Occurrence, Fate, and Effects of Pharmaceutical Substances in the Environment



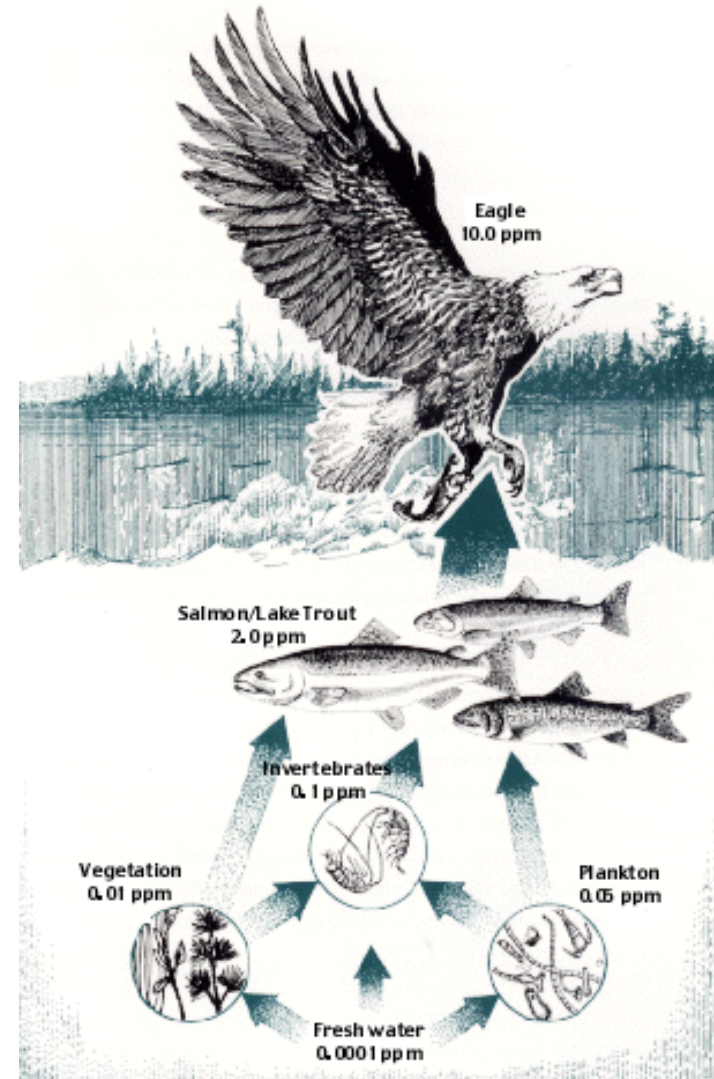
different exposure routes of veterinary and human medicinal substances

Partitioning:

K_{OW} , Indicator for Bioaccumulation

	$\log K_{OW}$
Benzene	2.13
Trichlorobenzene	4.05
Pentachlorophenol	5.01
PAHs	3.35 – 6.6
PCBs	~5 - ~7

$$K_{OW} = \frac{C_{oct}}{C_w}$$



Parasiticide Ivermectin:

Range of $\log K_{OW}$
3.2 (– 5.8) in the literature

Bioaccumulation?



Gaps in databases need to be filled!

Octanol-Water Partition Coefficient

Different Methods and Different Results

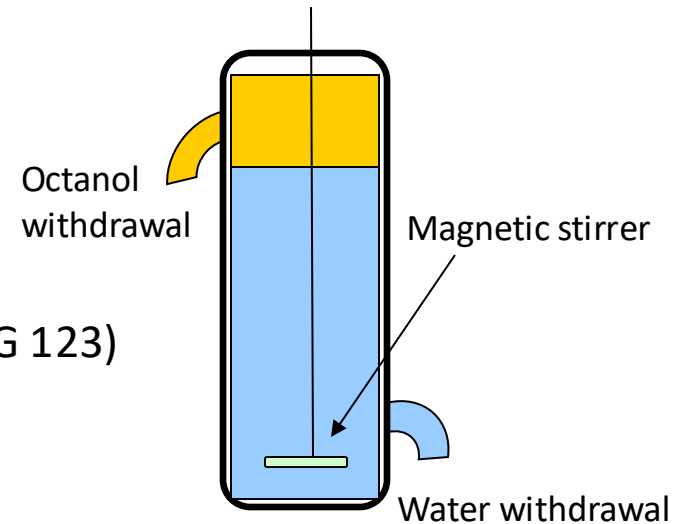


- Shaking method (OECD TG 107) with artifacts due to water droplets in octanol phase; up to $\log K_{ow}$ 4

- Slow stirring method (OECD TG 123) up to $\log K_{ow}$ 8



Correction of $\log K_{ow}$ for ivermectin:
from 3.2 to 5.6



(Römbke et al., 2019)

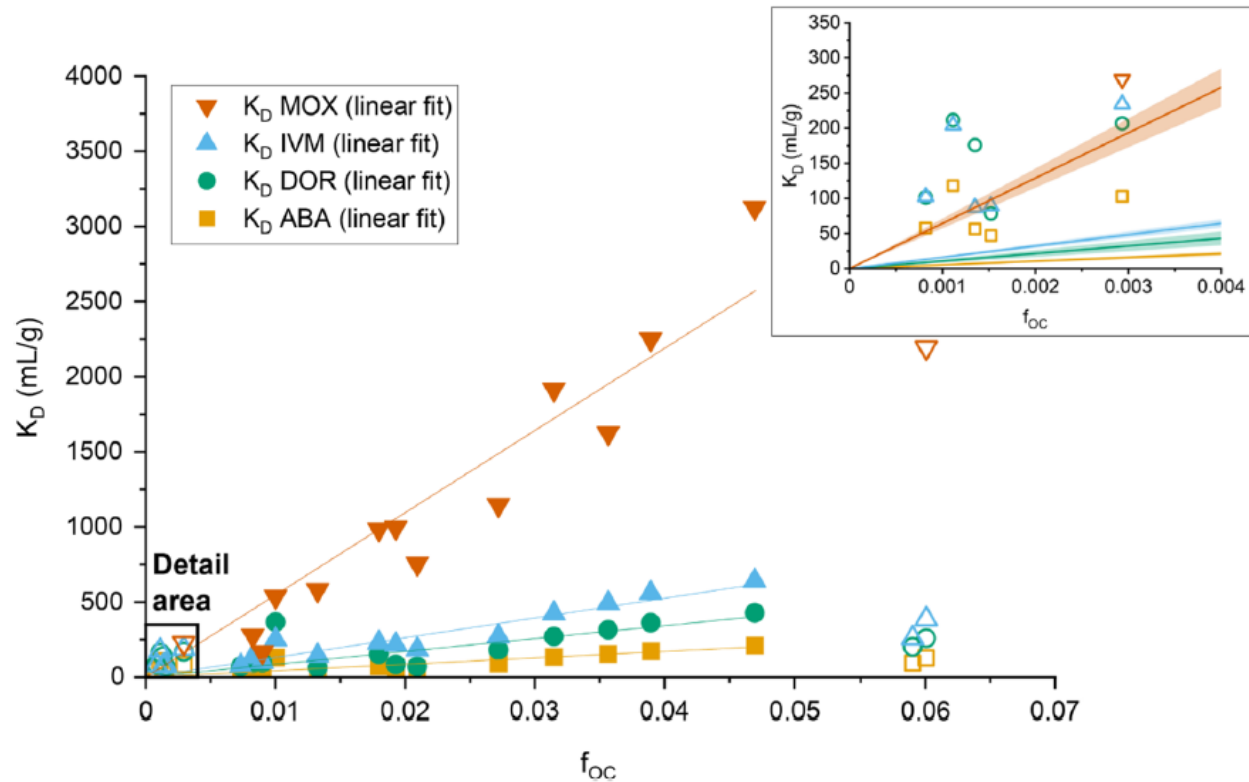
Sorption in Soils and Sediments

Indicator for mobility in the Environment

$$K_D = \frac{C_s(\text{eq})}{C_{\text{aq}}(\text{eq})}$$

$$K_{\text{OC}} = \frac{K_D}{f_{\text{OC}}}$$

K_{OC} is important for the classification of mobile substance as so-called „PMT“-substances



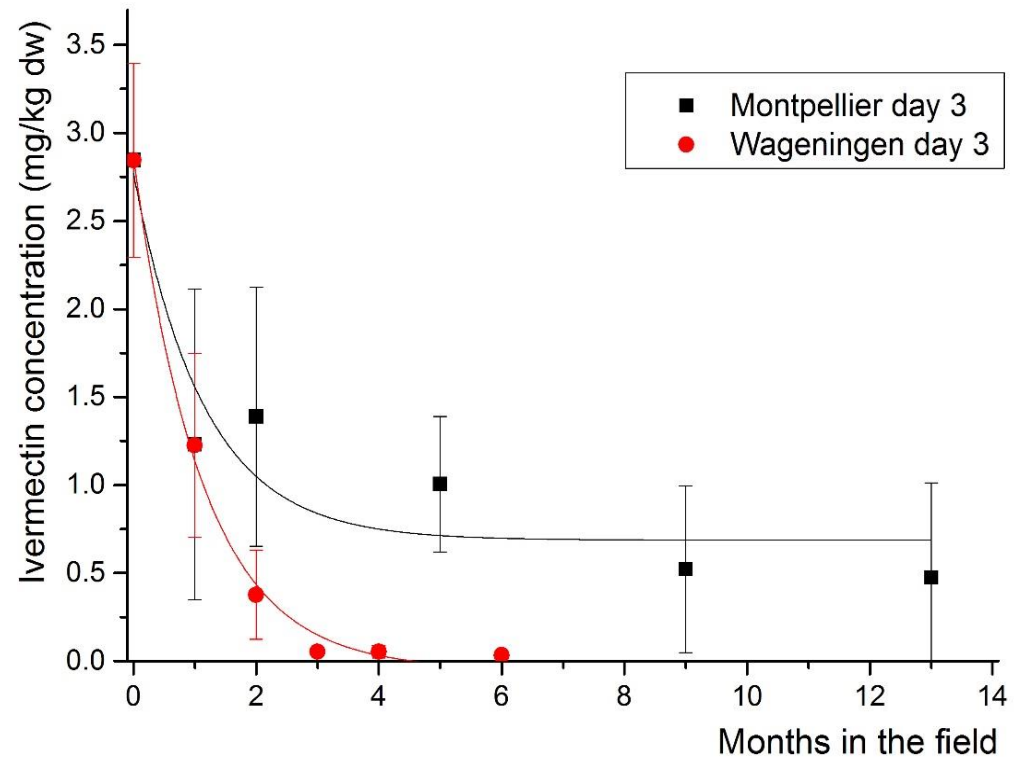
(Heinrich et al., 2021)

Transformation: Dissappearance, "Loss" of Ivermectin

Possible sub processes

- degradation?
- volatilization?
- leaching to surface and subsurface water?
- irreversible fixation to soil?
 - Transformation to non detectable compounds?

„...differences in ivermectin dissipation in cattle dung among sites, with 50% dissipation times of up to 32 d and 90% dissipation times >396 d.”



Wohde et al., 2016

Occurrence and Exposure

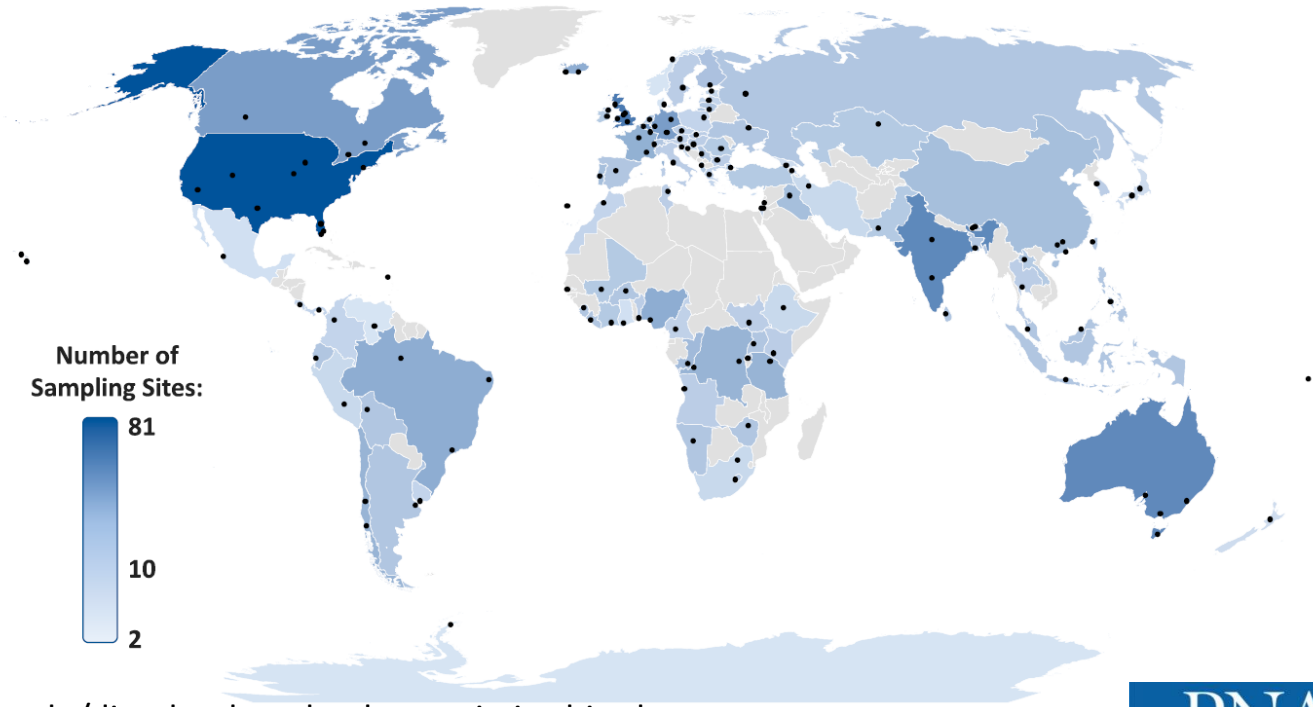
Pharmaceutical residues occur globally in the environment



(UBA, 2023)

Pharmaceutical pollution of the world's rivers

“...a global-scale study of API pollution in 258 of the world's rivers, representing the environmental influence of 471.4 million people across 137 geographic regions. Samples were obtained from 1,052 locations in 104 countries (representing all continents and 36 countries not previously studied for API contamination) and analyzed for 61 APIs...”



Databases:

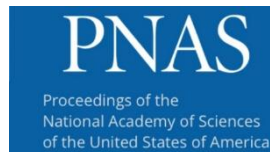
PHARMS-UBA

<https://www.umweltbundesamt.de/die-uba-datenbank-arzneimittel-in-der>

IPCHEM-Portal

<https://ipchem.jrc.ec.europa.eu/>

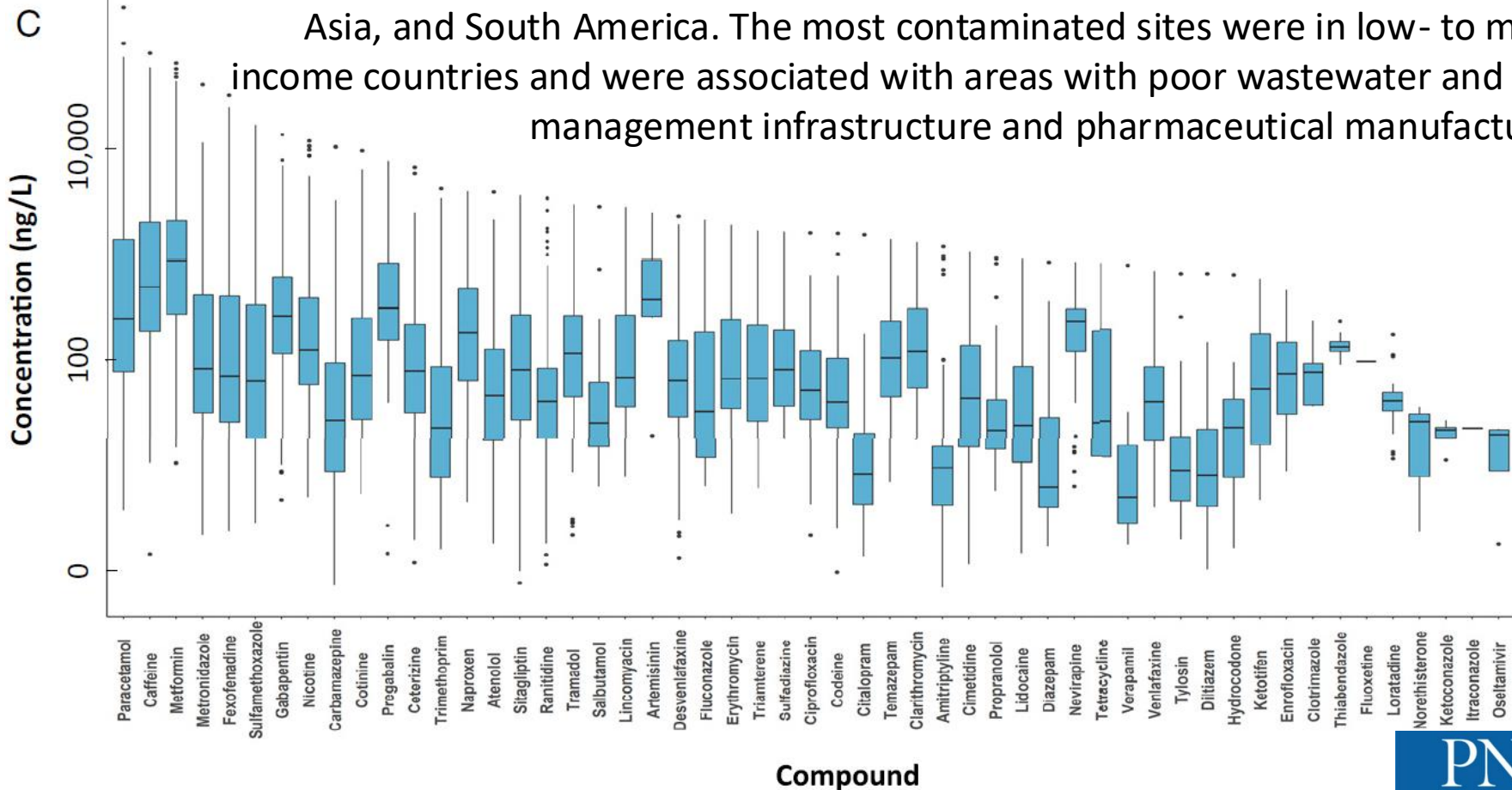
Wilkinson et al., 2022



Gießen, 03/04/2023

Pharmaceutical pollution of the world's rivers

“Highest cumulative API concentrations were observed in sub-Saharan Africa, south Asia, and South America. The most contaminated sites were in low- to middle-income countries and were associated with areas with poor wastewater and waste management infrastructure and pharmaceutical manufacturing.”



Wilkinson et al., 2022

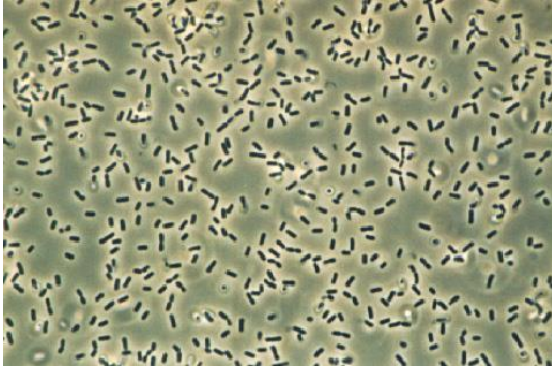
APIs and metabolites/transformation products in environmental compartments

Number of Year	Global		European Union	
	2021	Change to 2015	2021	Change to 2015
detected substances	992	+221	749	+153
in WWTP ¹ effluent/sewage/reclaimed water	771	+ 158	591	+117
in surface water/bank filtrate/groundwater/drinking and tap water	703	+ 175	483	+99
in manure/dung/sediment from aquaculture/SPM/biosolids/sludge	337	+192	250	+166
in sediment/soil/SPM	295	+111	227	+95

Source: UBA Pharms Database version 2 (2015) & version 3 (2021)

Gildemeister et al., 2022

Effects on Non-Target Organisms: Test Methods



- Inhibition of respiration of bacteria (Effect concentration, EC)
- Inhibition of reproduction of algae (Effect concentration, EC)
- Survival of crustacean (daphnie)
- Survival of fish (lethal dose, lethal concentration, LD/LC)

Studies on deputy organisms,
acute toxicity in mostly single-
species tests, standardized in
OECD Guidelines



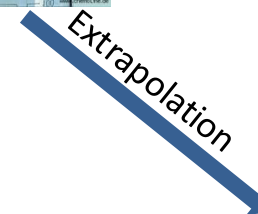
Effects of

pharmaceuticals on non-target organisms

Non-target organisms		Effect in Lab trial	Active substance
	water fleas	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #f08080; border: 1px solid #ccc;"></div> low toxic effect <div style="width: 20px; height: 10px; background-color: #800000; border: 1px solid #ccc;"></div> strong toxic effect </div>	Sulfadimethoxin, Sulfamethoxazol, Sulfadimidin, Trimethoprim Closantel, Cypermethrin, Deltamethrin, Doramectin, Eprinomectin, Fenbendazol, Flubendazol
	chironomidae	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #800000; border: 1px solid #ccc;"></div> strong toxic effect </div>	Deltamethrin
	fish	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #800000; border: 1px solid #ccc;"></div> strong toxic effect </div>	Altrenogest, Closantel, Cypermethrin, Deltamethrin, Eprinomectin, Ivermectin
	Earth worms	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #e91e63; border: 1px solid #ccc;"></div> moderate toxic effect </div>	Closantel, Cypermethrin, Deltamethrin, Eprinomectin, Ivermectin
	Dung organisms	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #e91e63; border: 1px solid #ccc;"></div> moderate toxic effect <div style="width: 20px; height: 10px; background-color: #800000; border: 1px solid #ccc;"></div> strong toxic effect </div>	Closantel Cypermethrin, Deltamethrin, Doramectin, Eprinomectin, Ivermectin
	soil organisms	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #ffc107; border: 1px solid #ccc;"></div> decreased phosphatase activity <div style="width: 20px; height: 10px; background-color: #ffc107; border: 1px solid #ccc;"></div> change of bacterial community </div>	Doxyzyklin Lincomycin, Sulfadiazin
	water plants	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #add8e6; border: 1px solid #ccc;"></div> low growth inhibition <div style="width: 20px; height: 10px; background-color: #0056b3; border: 1px solid #ccc;"></div> strong growth inhibition </div>	Trimethoprim Florfenicol
	crops	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #007bff; border: 1px solid #ccc;"></div> moderate germination inhibition <div style="width: 20px; height: 10px; background-color: #0056b3; border: 1px solid #ccc;"></div> strong germination inhibition <div style="width: 20px; height: 10px; background-color: #007bff; border: 1px solid #ccc;"></div> moderate germination inhibition <div style="width: 20px; height: 10px; background-color: #0056b3; border: 1px solid #ccc;"></div> strong germination inhibition </div>	Sulfamethoxazol Florfenicol Enrofloxacin, Sulfadiazin Enrofloxacin, Florfenicol
	cyano bacteria	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #add8e6; border: 1px solid #ccc;"></div> low growth inhibition <div style="width: 20px; height: 10px; background-color: #007bff; border: 1px solid #ccc;"></div> moderate growth inhibition <div style="width: 20px; height: 10px; background-color: #0056b3; border: 1px solid #ccc;"></div> strong growth inhibition </div>	Trimethoprim Amoxicillin/Penicillin Säure, Tetrazyklin Enrofloxacin, Erythromycin, Oxytetrazyklin
	green algae	<div style="display: flex; justify-content: space-between;"> <div style="width: 20px; height: 10px; background-color: #ccc; border: 1px solid #ccc;"></div> no growth inhibition <div style="width: 20px; height: 10px; background-color: #007bff; border: 1px solid #ccc;"></div> moderate growth inhibition <div style="width: 20px; height: 10px; background-color: #0056b3; border: 1px solid #ccc;"></div> strong growth inhibition </div>	Amoxicillin/Penicillin Säure Enrofloxacin, Ivermectin, Tetrazyklin Erythromycin



Laboratory



Ecosystem



toxic effect

shift of species composition

growth inhibition

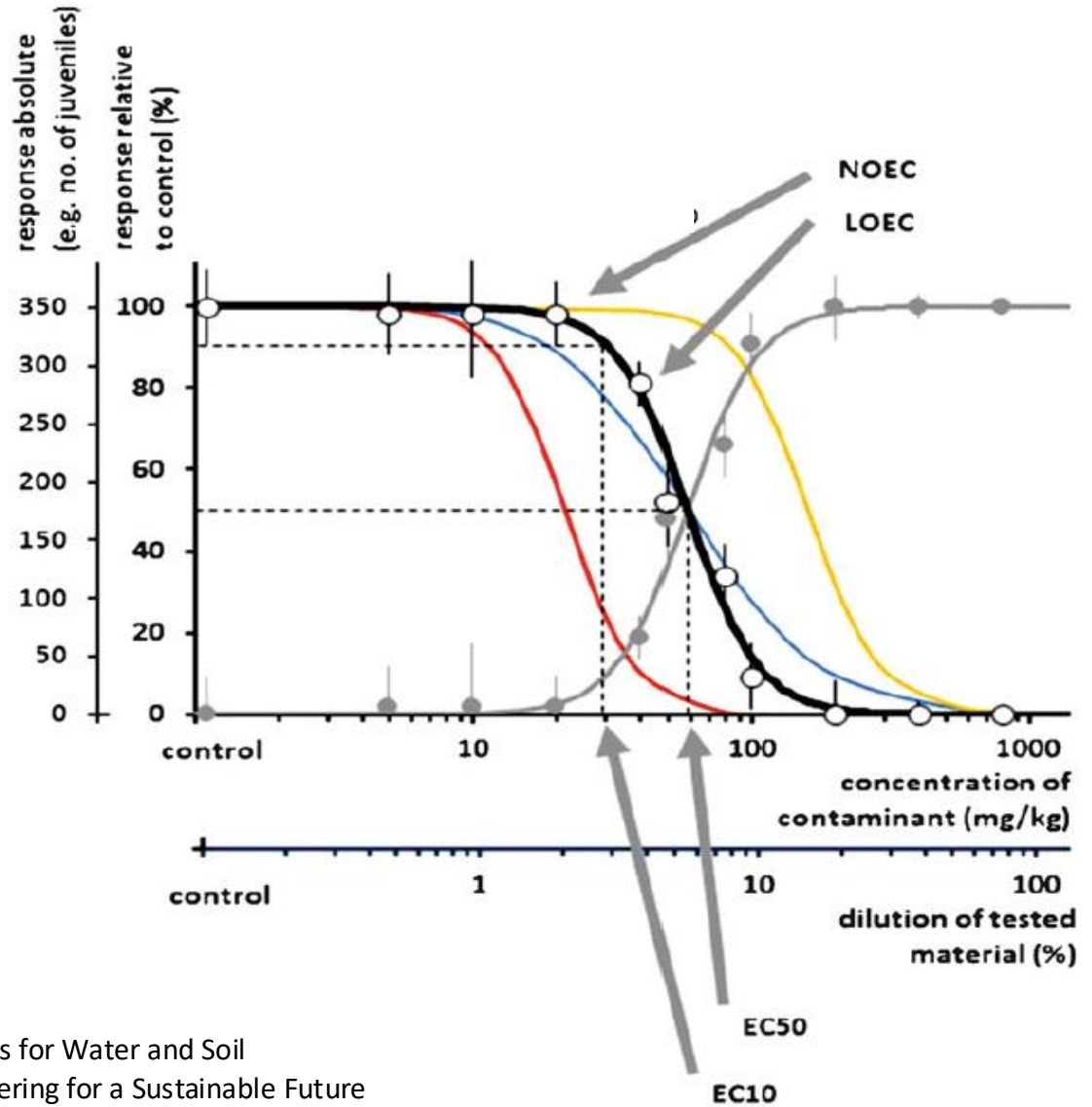
(UBA, modified, 2017)

Gießen, 03/04/2023

Prospective Assessment: Ecotoxicity of Chemicals

Ecotoxicity parameters

(**NOEC**, **LOEC**, **EC50**, **LC50**, etc.)
are derived from the modelled
concentration–response curve,
determined for one single test
substance

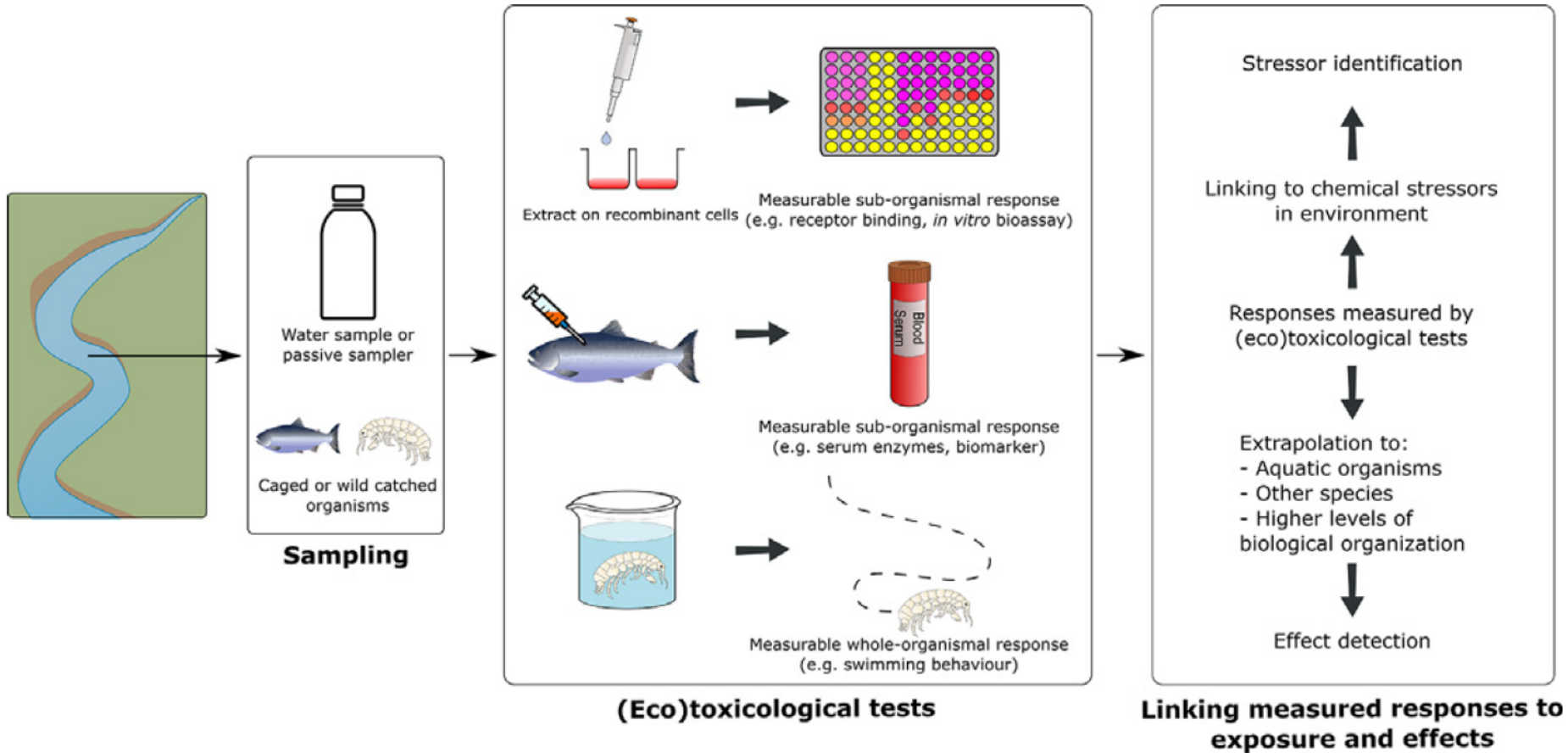


Adapted from Bláha and Hofman, 2020

in: J. Filip et al. (eds.), Advanced Nano-Bio Technologies for Water and Soil
Treatment, Applied Environmental Science and Engineering for a Sustainable Future

Retrospective Assessment: Ecotoxicology of Contaminated Samples

Advanced approach: integration of ecotoxicity tests into monitoring practices



Schuijt et al. (2021)

Environmental Risk Assessment Based on Exposure and Effect



danger: lion



exposure: low in Europe

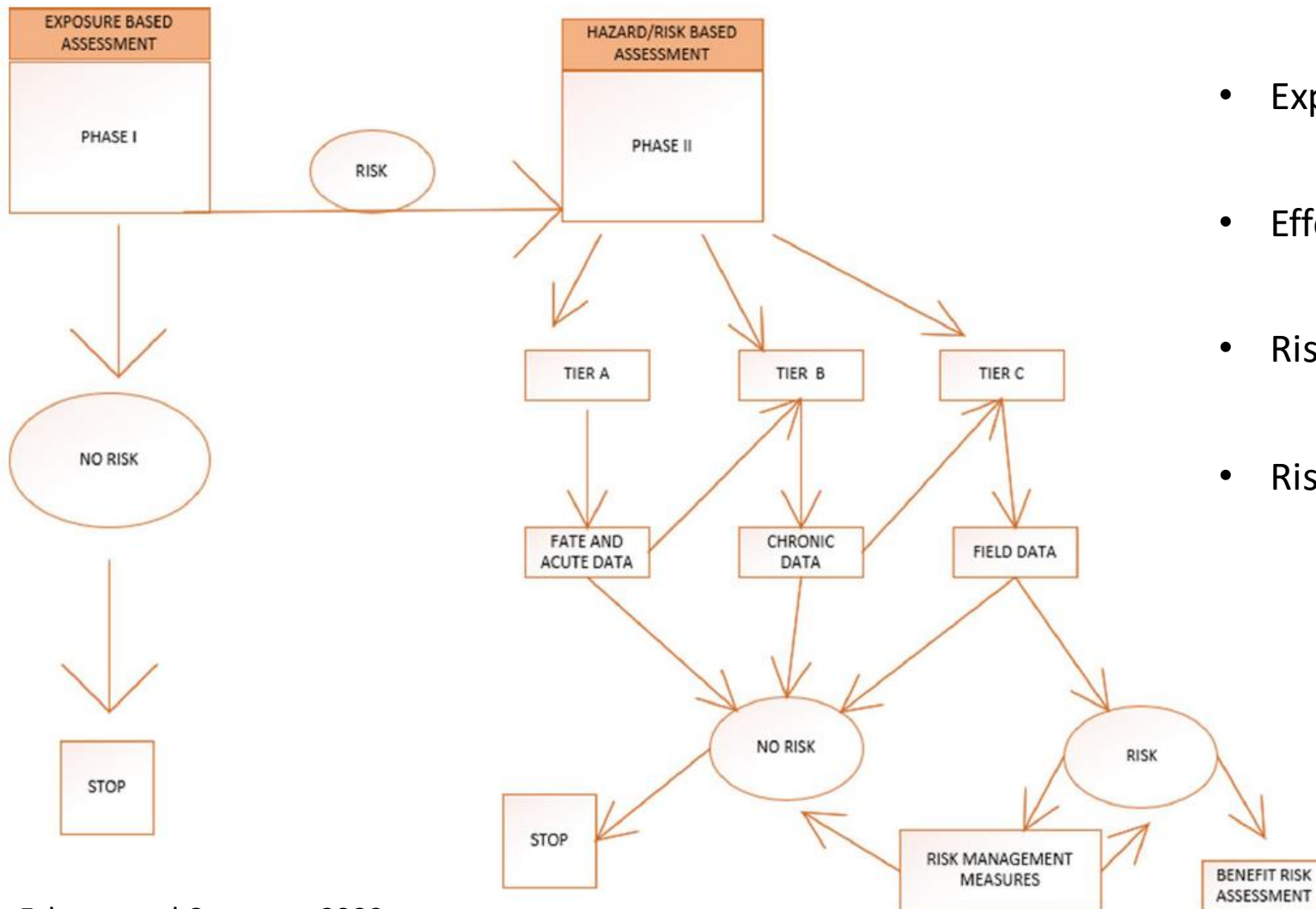
“danger“: corresponds to **PNEC**,
predicted no effect
concentration;
e.g. on the basis of NOEC
including a safety factor

“probability of exposure“:
corresponds to **PEC**, predicted
environmental concentration;
sometimes “MEC“ is used
instead

Environmental risk as a result of intrinsic hazard
of a compound and probability for exposure

The aim is:
$$\frac{PEC}{PNEC} < 1$$

Framework for the Environmental Risk Assessment for Veterinary Medicinal Products in the EU



- Exposure
- Effects
- Risk Assessment
- Risk Management

Fabrega and Carapeto, 2020

Environmental Risk Assessment (ERA)

International Governance

- ERA principles for veterinary pharmaceuticals are defined in an international framework (VICH)
- Two basic guidelines: VICH GL 6 & VICH GL 38
- Step-wise approach: Phase I and Phase II ERA

VICH GL6 (ECOTOXICITY PHASE I)
June 2000
For implementation at Step 7

**ENVIRONMENTAL IMPACT
ASSESSMENT
(EIAs) FOR VETERINARY MEDICINAL
PRODUCTS (VMPs) - PHASE I**

Recommended for Implementation
at Step 7 of the VICH Process
on 15 June 2000
by the VICH Steering Committee

THIS GUIDELINE HAS BEEN DEVELOPED BY THE APPROPRIATE VICH EXPERT WORKING GROUP AND WAS SUBJECT TO CONSULTATION BY THE PARTIES, IN ACCORDANCE WITH THE VICH PROCESS. AT STEP 7 OF THE PROCESS THE FINAL DRAFT IS RECOMMENDED FOR ADOPTION TO THE REGULATORY BODIES OF THE EUROPEAN UNION, JAPAN AND USA.

**ENVIRONMENTAL IMPACT
ASSESSMENT FOR VETERINARY
MEDICINAL PRODUCTS
PHASE II GUIDANCE**

Recommended for Adoption
at Step 7 of the VICH Process
in October 2004 by the VICH SC for implementation in October 2005

This Guidance has been developed by the appropriate VICH Expert Working Group and is subject to consultation by the parties, in accordance with the VICH Process. At Step 7 of the Process the final draft will be recommended for adoption to the regulatory bodies of the European Union, Japan and USA.

Phase I ERA

Determination of **environmental exposure of the medicine** and need for ecotoxicological assessment, **mandatory for all veterinary medicines.**

Based on exposure

- Low exposure → Limited risk
- Individual treatments → Phase I
- Pets → Phase I
- Natural substances → Phase I
- Exposure < 100 µg/kg → Phase I

Specific issues:

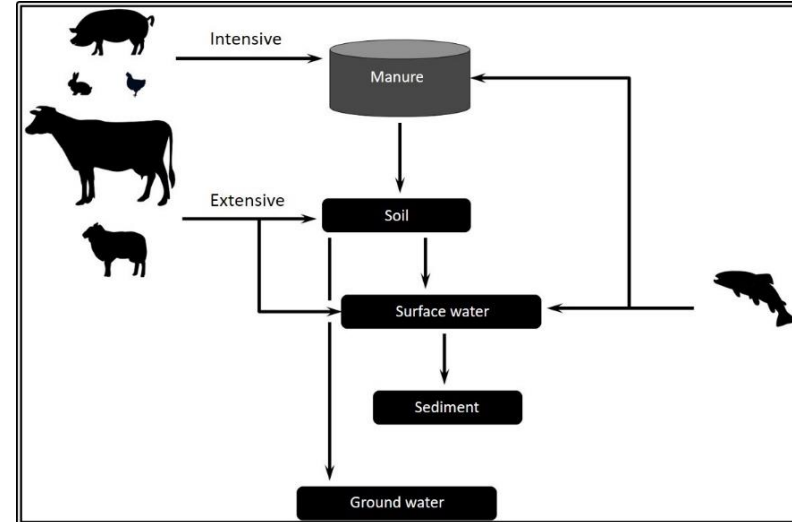
- Parasiticides → Phase II
- Aquaculture open waters → Phase II

- “However clause”

Relevant for majority of pharmaceutical veterinary medicinal products (> 95%)

Phase II ERA

- Problem formulation – Protection goals
 - Protection of ecosystems



- Risk assessment

- Exposure calculation (PEC)



Husbandry practices

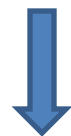
Physicochemical/fate studies

- Toxicity determination (PNEC)



Environmental compartments

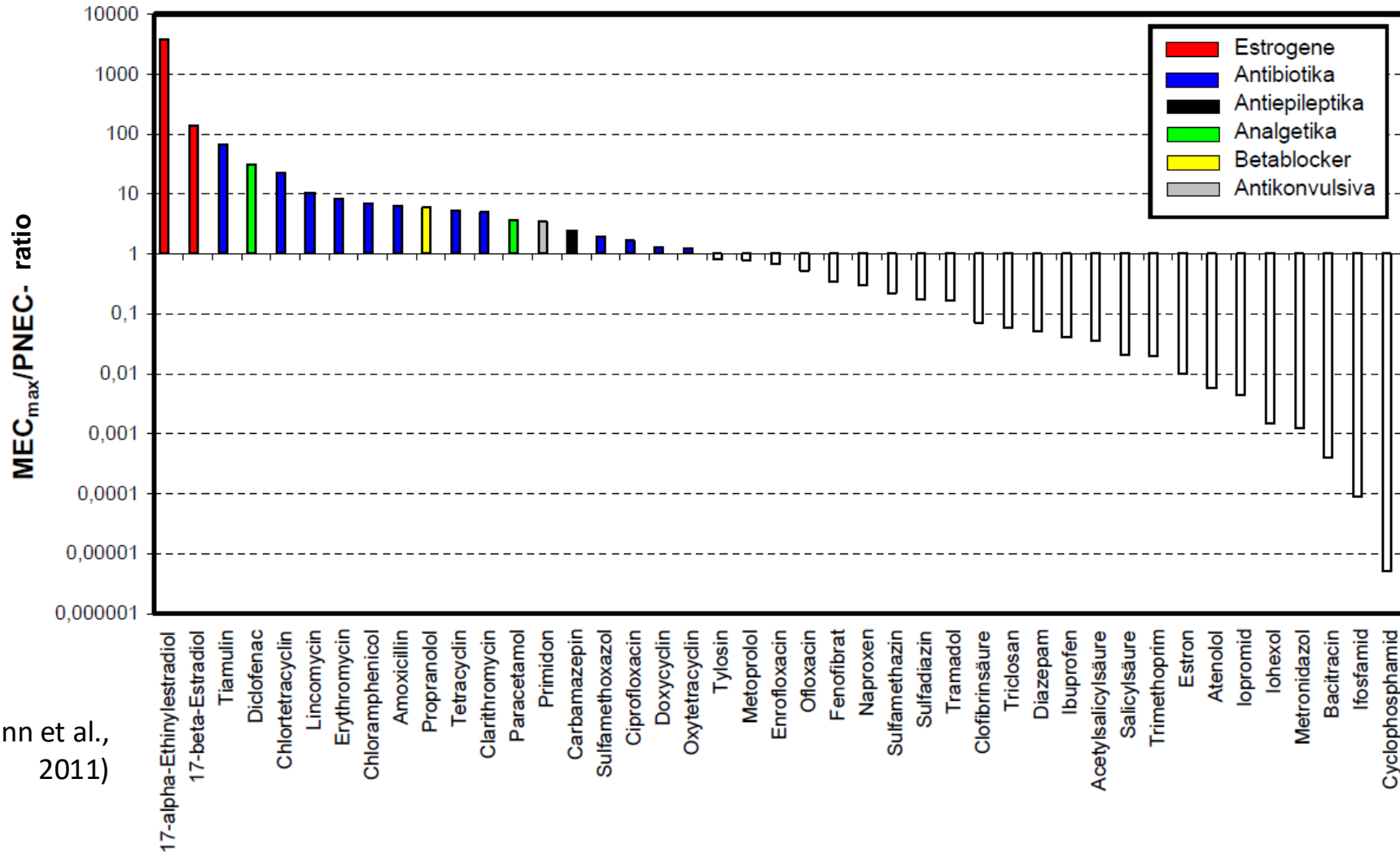
- Soil
- Water
- Dung
- Sediment



- Risk Quotient Approach $\rightarrow RQ = PEC/PNEC$

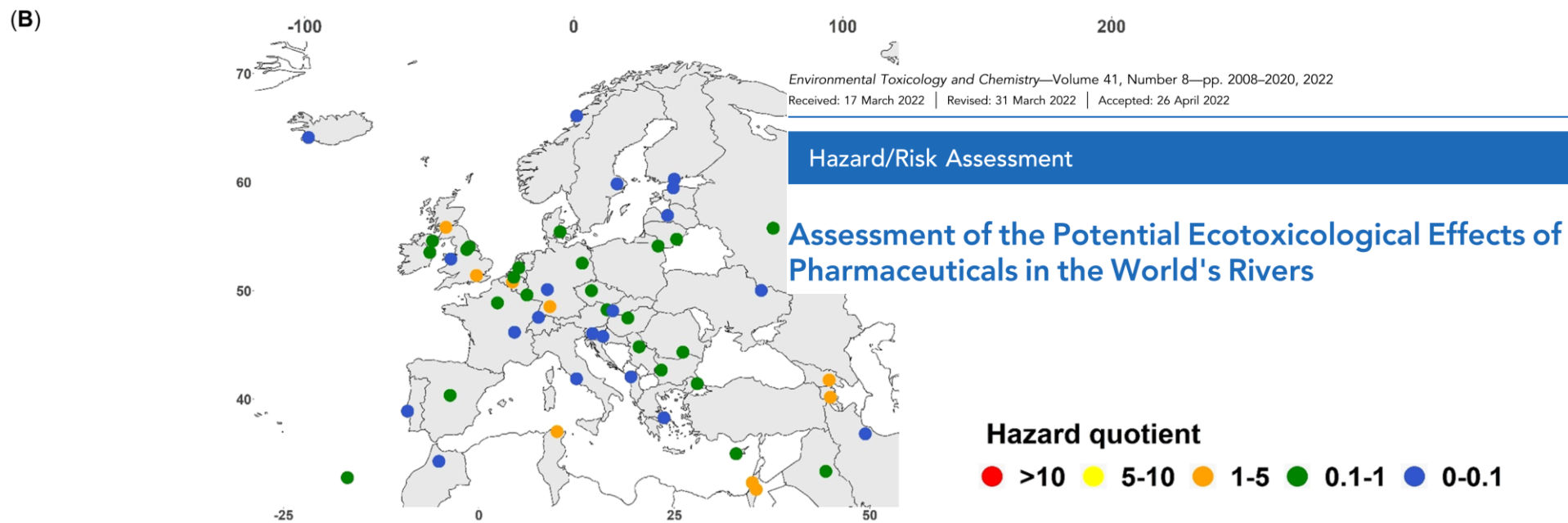
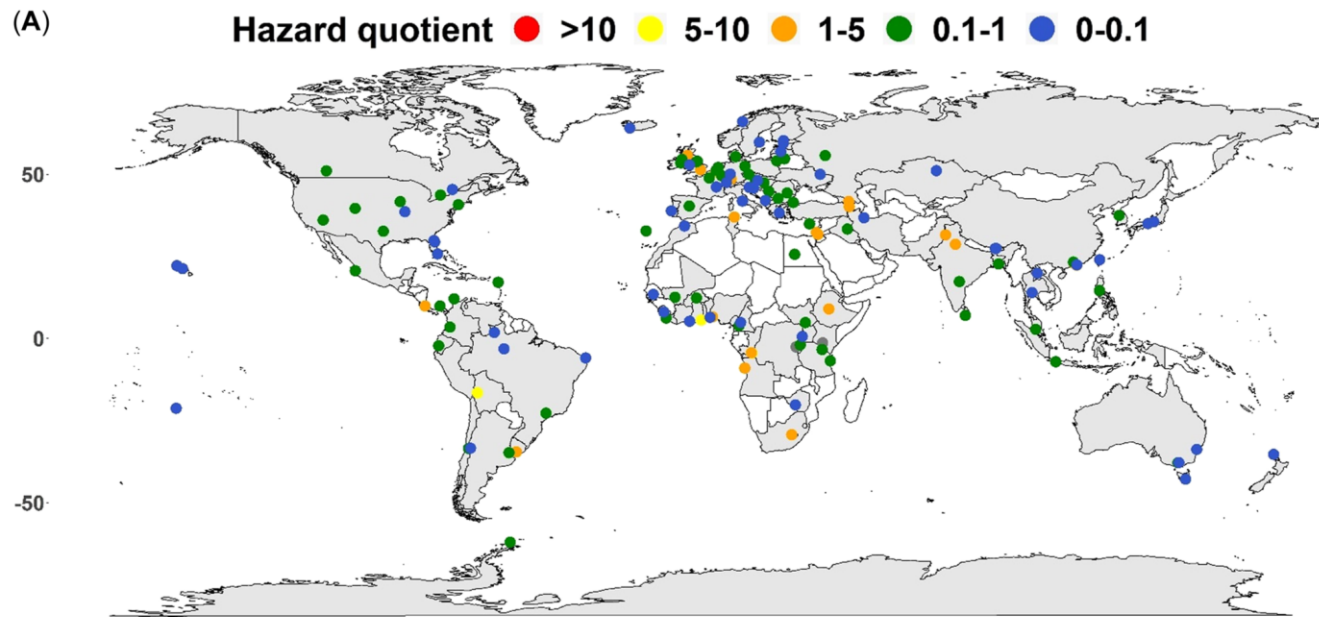
mitigation measures to reduce the risk to an acceptable level?

MEC/PNEC-Ratios for Pharmaceuticals with Good to Sufficient Ecotoxicological Data Base



(Bergmann et al.,
2011)

Risk Assessment



Little Consideration of APIs in Environmental EU legislation

watchlist:

- diclofenac
- estrone (E1)
- 17-beta-estradiol (E2)
- 17-alpha-ethinylestradiol (EE2)
- three macrolide antibiotics

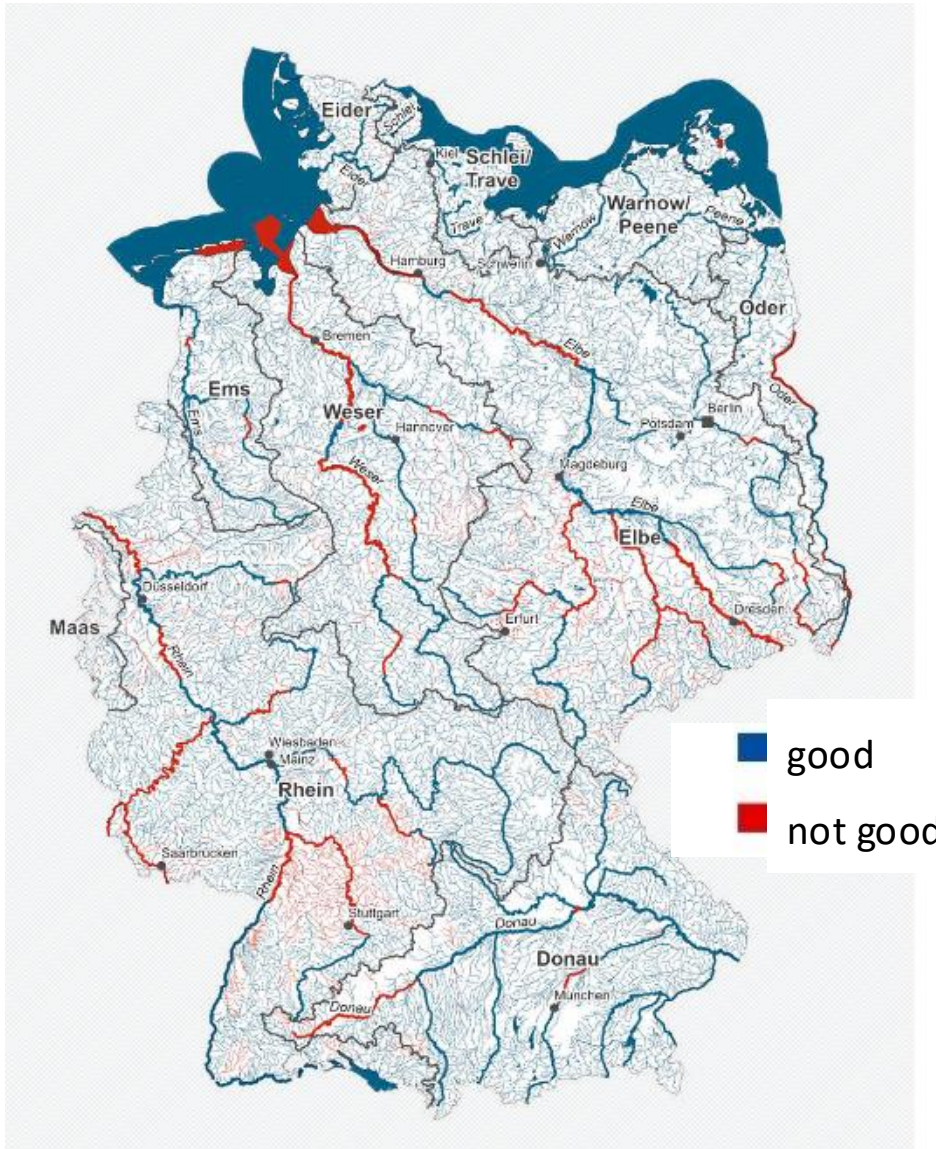
EU Legislative framework	Year of adoption	Consideration of pesticides ¹⁾	Consideration of pharmaceuticals	Current EU action
Water Framework Directive (2000/60/EC)	2000	Annex VI, Part A Annex VIII	No	-
Directive on Environmental Quality Standards (Directive 2008/105/EC amended by 2013/39/EU)	2013	Art. 7a, Substances in annex (EQS)	Art 8b Watchlist, Art 8c strategic approach	Review of priority substances ²⁾ , Fulfilling strategic approach pharmaceuticals ³⁹
Groundwater Directive (2006/118/EC)	2006	Annex I: Groundwater quality standards	No	-
Sewage Sludge Directive	1986	No	No	revision proposed, impact assessment closed
Urban Waste Water Directive	1991	No	No	in revision
Industry Emissions Directive	2010	Annex I: Chemical industry / production	Annex I: Chemical industry / production	draft of revision published in 2022
Classification, labelling & packaging regulation	2008	YES	No	Revision planned
Revised Drinking Water Directive	2020	(17) Recital Annex I Water quality	(7), (17) Recital Art. 13 8. Monitoring Art. 19 3. Evaluation	-
Soil Health Law	Open	Open	Open	Proposal in 2023

Gildemeister et al., 2022

1) regulations on plant protection products and biocides

2) watch list (established in 2015, updates: 2018, 2020, 2022): pharmaceuticals substances included and now proposed as candidates for priority substances

Water Framework Directive Chemical Status of Surface Waters (ubiquitous pollutants excluded)

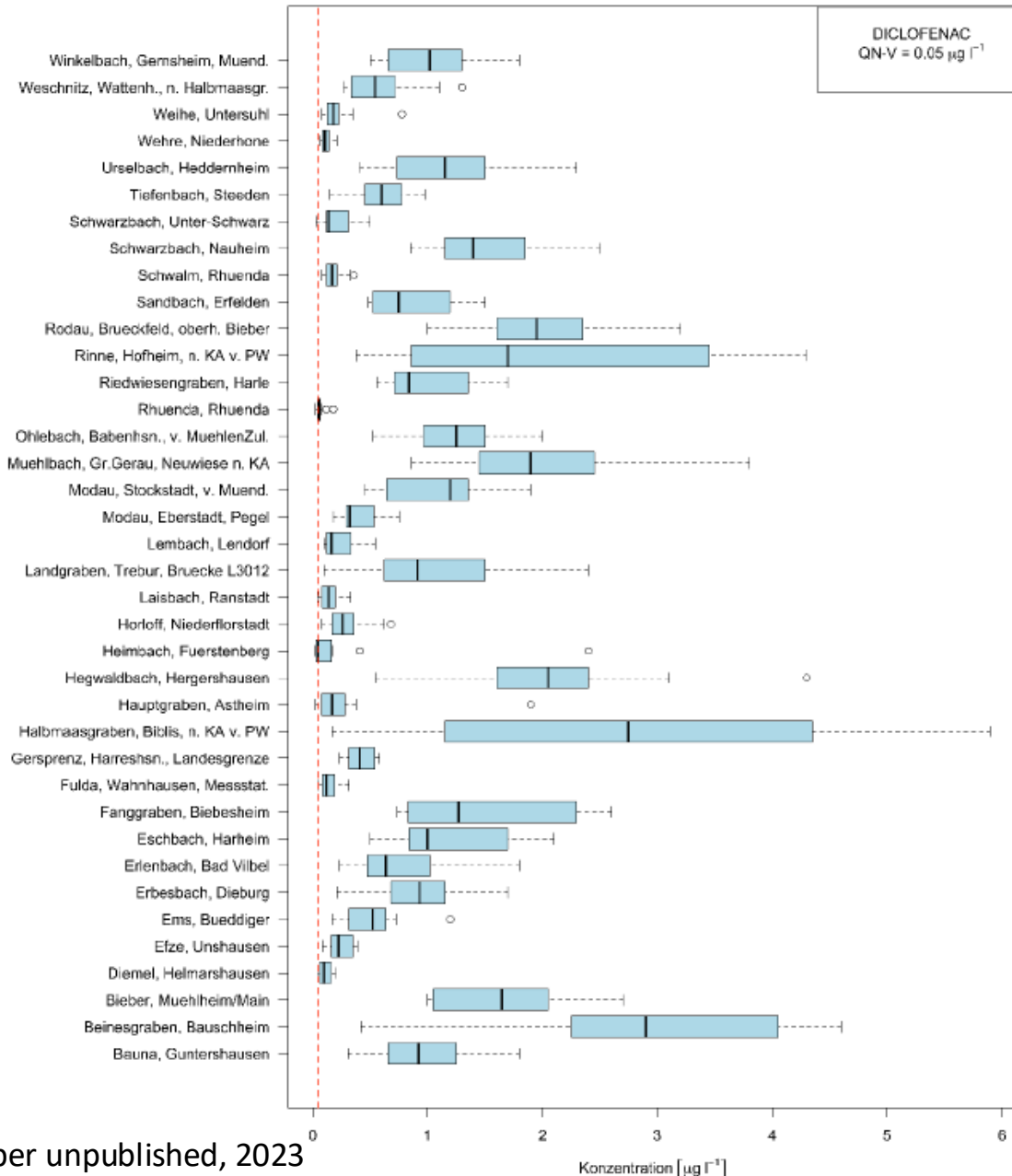


the aim is to bring all rivers, lakes, groundwater and coastal waters into a "good status" by 2027 at the latest.

assessment is based on concentrations in water and biota

Pharmaceuticals in Running Waters in Hesse, Germany

Diclofenac

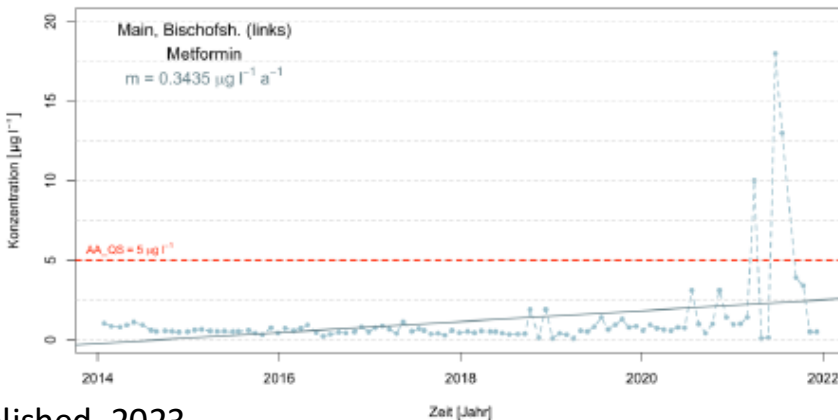
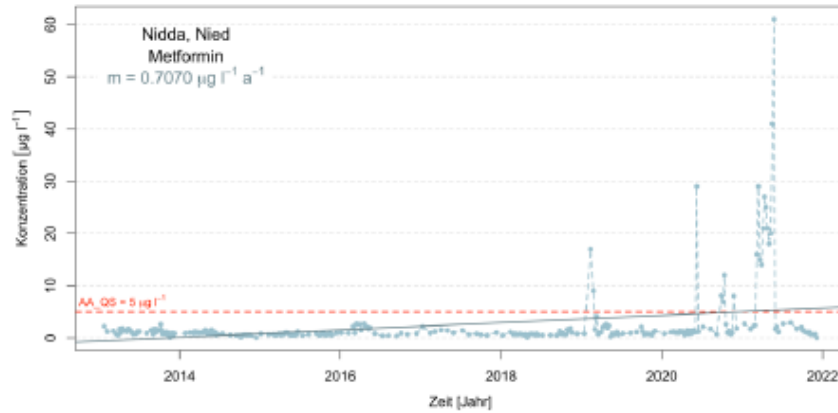
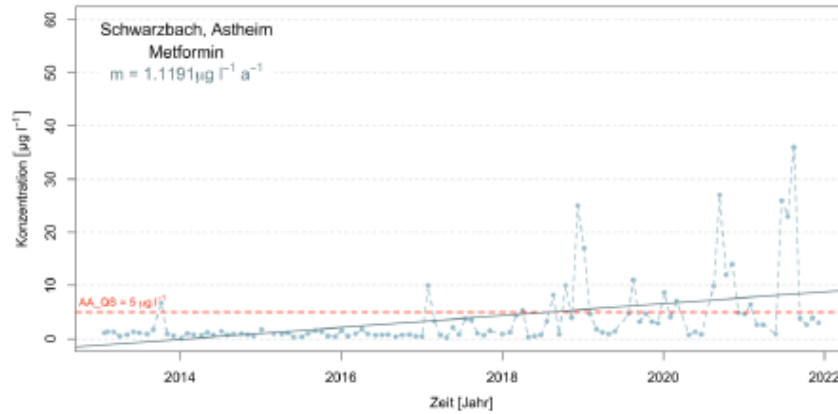


Always above quality limit (quality standard proposal), with increasing tendency!

This would mean every river with “not good chemical quality”

Pharmaceuticals in Running Waters in Hesse, Germany

Metformin



Recent strong upward trend
(transformation product not considered)!

Solutions for Sustainable Management(?)

our Basis for Discussion

Drug design

- Publicly accessible collection of data
- Strengthening of ERA with the possibility of refusal of approval
- Research promotion of green pharmacy
- Introduction of an environmental classification system for pharmaceuticals
- Extension of good manufacturing practice to include environmental requirements

Application

- Expansion of preventive and precautionary health care
- Information and training of healthcare professionals
- Prescription requirement for drugs that are particularly hazardous to the environment
- Prohibition of advertising for non-prescription drugs

Downstream measures

- Centralized collection of pharmaceutical residues nationwide via the pharmacies
- Expansion of the 4th purification stage primarily at polluted wastewater treatment plants; separation hot spots (e.g. hospitals) or X-ray contrast media by separate collection and disposal collection and disposal

Costs

- According to the polluter pays principle, pharmaceutical manufacturers should contribute to the reduction measures

Thank you for listening!

