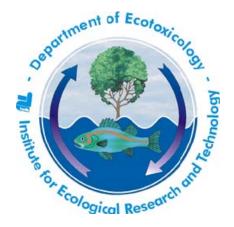
# Risk Assessment in Aquatic Systems

#### Peter D. Hansen

Technische Universität Berlin, Institute for Ecological Research and Technology, Department of Ecotoxicology, Franklinstrasse 29 (OE4), D-10587 Berlin, Germany



www.tu-berlin.de/~oekotox www.ecosystemhealth.net

#### **Overview**

#### Risk

- Assessment
- Communication
- Managemnet

**Exposure Toxicity Ratio ETR Szenarios: sediment / soil / water** 

**Quality Standards: EQS** 

**Quality Norms: QN, UQN, EQN** 

**Ecological Risk Assessment** 

**Roadway - Implementation WFD** 

The good ecological status

The good chemical status

Effect related Bioassays are in use for the "good chemical status": Quality Norms (QN – [μg/L])

**REACH** and surface water protection

**Ecological Risk Assessment** 

#### **Risk Assessment**

#### **Risk Definition:**

**Proparibility and Quantification of Disruptions by Hazards** 

#### **Process of Risk Assessment:**

- 1. Identification of Risk
- 2. Effects Assessment
- 3. Exposure Assessment
- 4. Risk Characterisation

# Concept of the Risk Assessment

#### **Exposure**

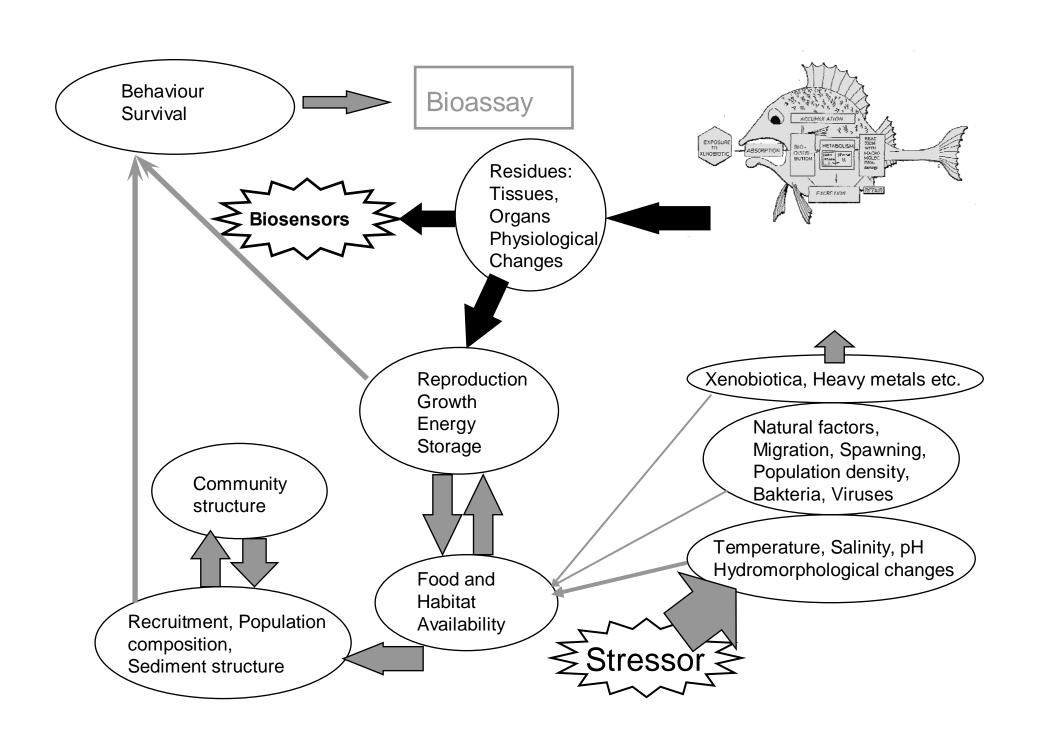
(Exposure Assessment)

- → Compounds Sale
- → Formulation and active ingredient
- → Comp. structural properties/effecs
- → Exposure matrices and routing
- → Biotransformation
- → Metabolites
- **→**Excretion
- → Kinetics and Excretion
- → Biological degradation (Persistence)
- → Bioavailability

#### **Effects**

(Effects Assessment)

- → Accumulation
- → acute and/or chronic Toxicity
- → Enzyme induction
- → Enzyme inhibition
- → Antibiotica Resistence
- → Reproductions Toxikology



# Effects Assessment - Environment and Human Health Environmental Signalling

1 Reaction



2 ) Adaptation



3 )

**Death** 

Biosensors Endpoints

→Enzyme-Induction Adap

- → Enzym-Inhibition
- → Biotransformation
- → Oxiradicals
- → Gentoxicity
- → Endocrine Effects
- **→**Immuntoxicity
- → Proteinbinding (Metals)

real time - Bioassays Large scale Bio-Monitoring Chronic Effects

#### **Adaptive Enzymes**

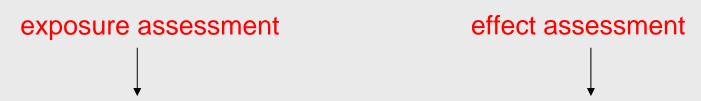
- →Inhibition of Functions
- → Disruption of Structures
- → Reconstitution

#### **Change of Behaviour**

- → Populations selection
- → Change in Reproductionstrategy

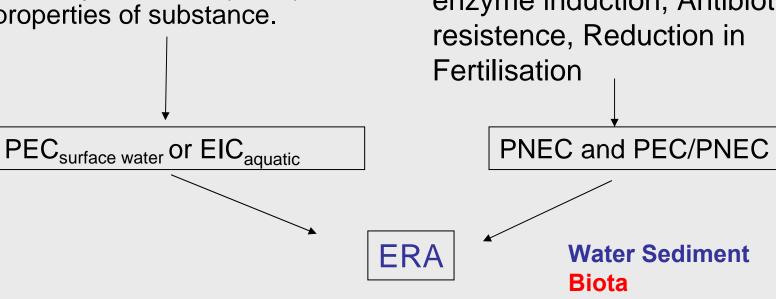
Lost of important Biochemical Functions

# **Environmental Risk Assessment**

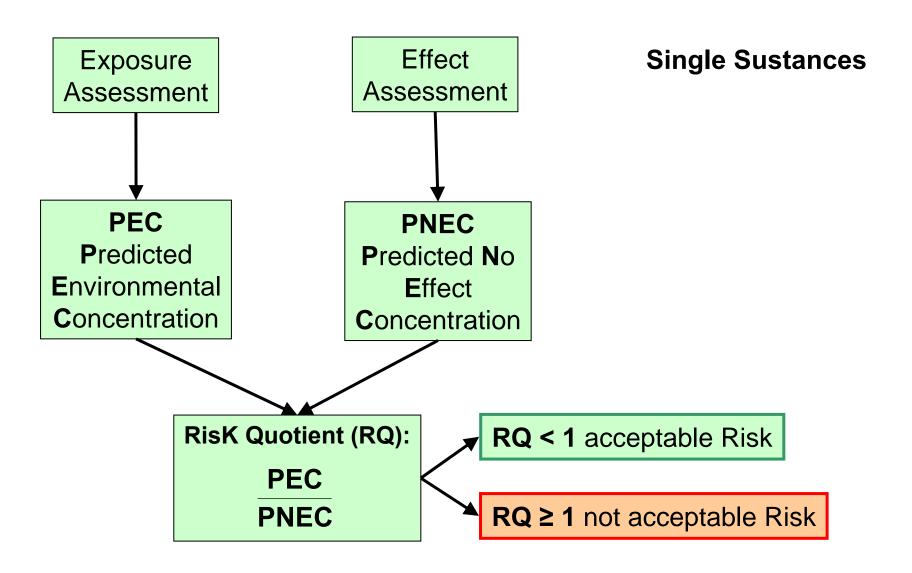


biotransformation; metabolites; exposure pathway; excretion; retention period; biodegradability; market penetration (sales); properties of substance.

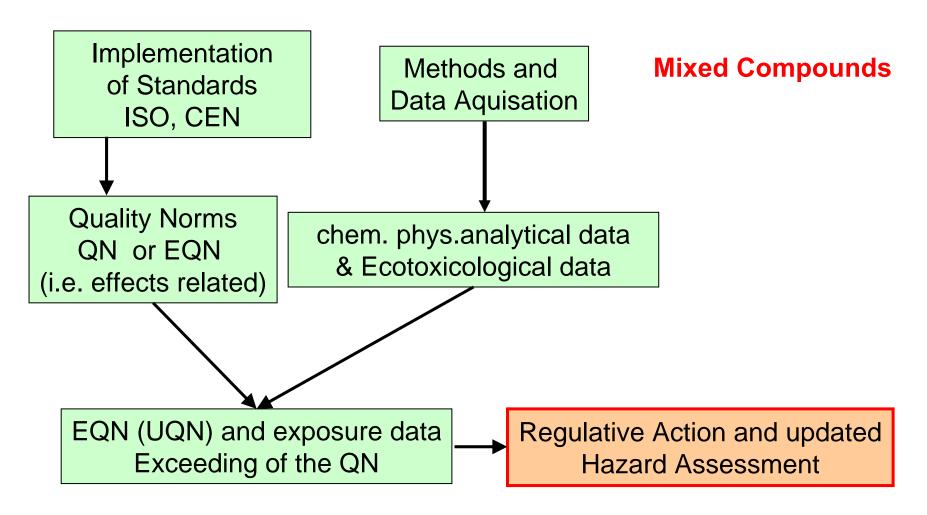
accumulation; acute und chronic effects; adverseor indirect effects; (e.g. enzyme induction, Antibiotica resistence, Reduction in Fertilisation

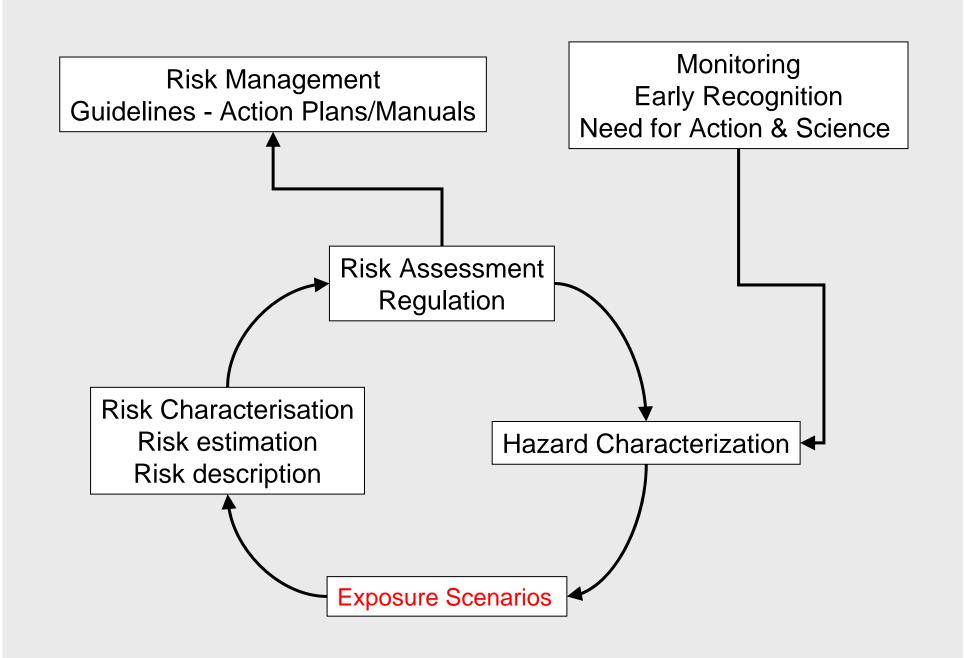


# Concept of the Risk Assessment

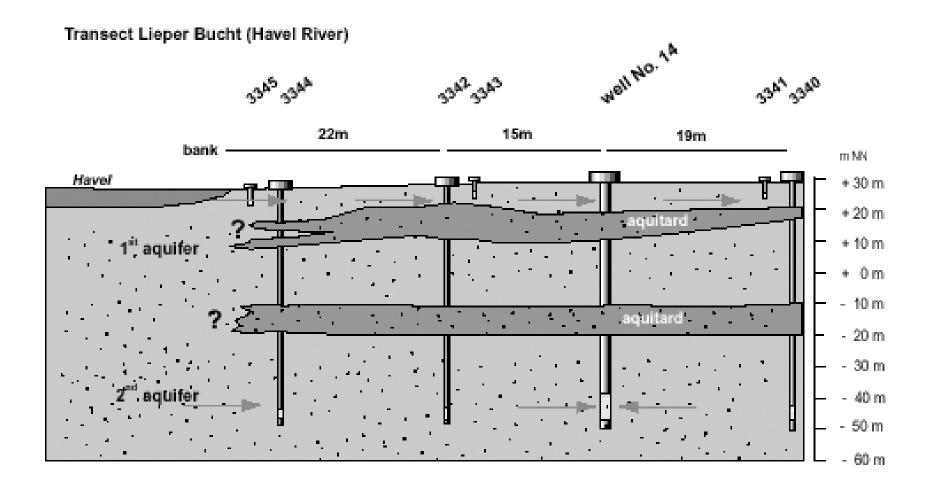


# Concept of Hazard Assessment









Enzelstoff	Konzentration in ng/l
2,4-D	ND- 20
Bentazone	ND - 50
Mecoprop	ND - 130
o,p´-DDA	ND - 25
p,p´-DDA	ND - 90
NPS*	15 -1180
TCEP**	20 - 270
TCIPP***	65 - 1160
Bezafibrate	ND - 420
Compound X	ND - 830
Carbamazepine	25 - 1075
Clofibric Acid	2 - 450
Caffeine	80 - 970
Diclofenac	ND - 1030

Einzelstoff	Konzentration in ng/l
Fenofibric Acid	up to 700
Gemfibrozil	ND - 85
Ibuprofen	ND - 75
Indomethacine	ND - 85
Ketoprofen	ND - 65
Mefenamic Acid	ND - 20
Naproxen	ND - 95
Oxazepam	ND - 85
Pentoxifylline	ND - 30
Primidone	ND - 635
Propiphenazone	ND- 1970
Tolfenamic Acid	ND - 20

Ethinylestradiol:	1.0 ng/l
17ß-Estradiol	0.5 ng/l
Estron	0.9 ng/l

Nonylphenol	110 ng/l
Diallylphtalat	225 ng/l
Bisphenol A	31 ng/l

<sup>\*</sup> N-(Phenylsulfonyl)-Sarcosine \*\* Tris-(ChloroEthyl)-Phosphate

<sup>\*\*\*</sup> Tris-(ChloroIsoPropyl)-Phosphate

# Composited samples in Sedimentlayers 0-5 cm

	4° C	27° C	pH 4	pH 10
Einwaage (g)	2,01	2,04	2,21	2,15
PAK in μg/g TS				
Naphthalen	4,4	1,4	2,4	1,5
Acenaphthylen	1,5	0,9	1,2	1,5
Acenaphthen	4,4	1,8	3,8	1,5 1,5 3,2 9,5
Fluoren	9,3	5,6	10,6	
Phenanthren	21,1	18,1	29,2	25,0
Anthracen	10,0	8,5	12,2	11,2
Fluoranthen	35,4	38,9	52,0	41,1
Pyren	31,4	32,5	44,7	37,1
Benz(a)anthracen	21,3	24,9	31;2	28,0
Chrysen/Triphenylen	21,1	23,5	30,0	27,3
Benzo(b,j)fluoranthen	21,3	23,8	29,1	27,5
Benzo(k)fluoranthen	14,1	15,7	18,9	18,2
Benzo(e)pyren	22,5	24,4	31,0	28,3
Benzo(a)pyren	21,3	24,3	27,9	27,6
Perylen	11,0	11,3	13,8	13,8
Dibenzo(a,h)anthracen	7,7	7,3	9,5	10,7
Indeno(1,2,3-cd)pyren	15,8	17,2	18,7	21,5
Benzo(g,h,i)perylen	16,3	17,1	18,6	22,2
Σ PAK (12)	236,9	251,9	322,0	297,4
ΣΡΑΚ	290,0	297,4	384,7	355,2

#### **Risk Assessment - Sediments**

(Eco) –toxic effect



Toxicity – Classification pT (EQN - UQN)



**Bioassays** 

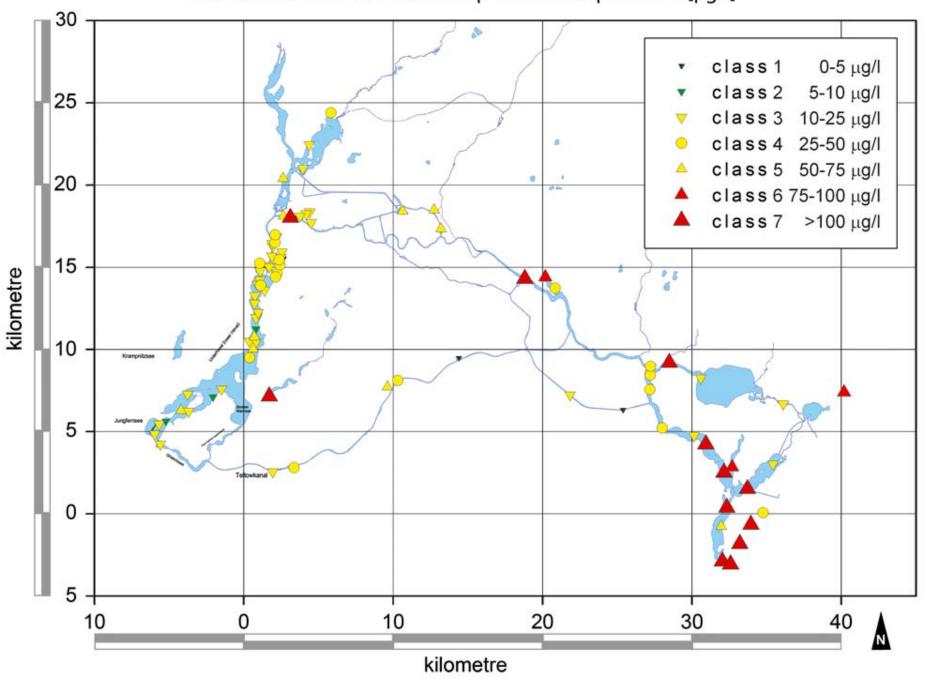


**Dilutionsteps (LID)** 



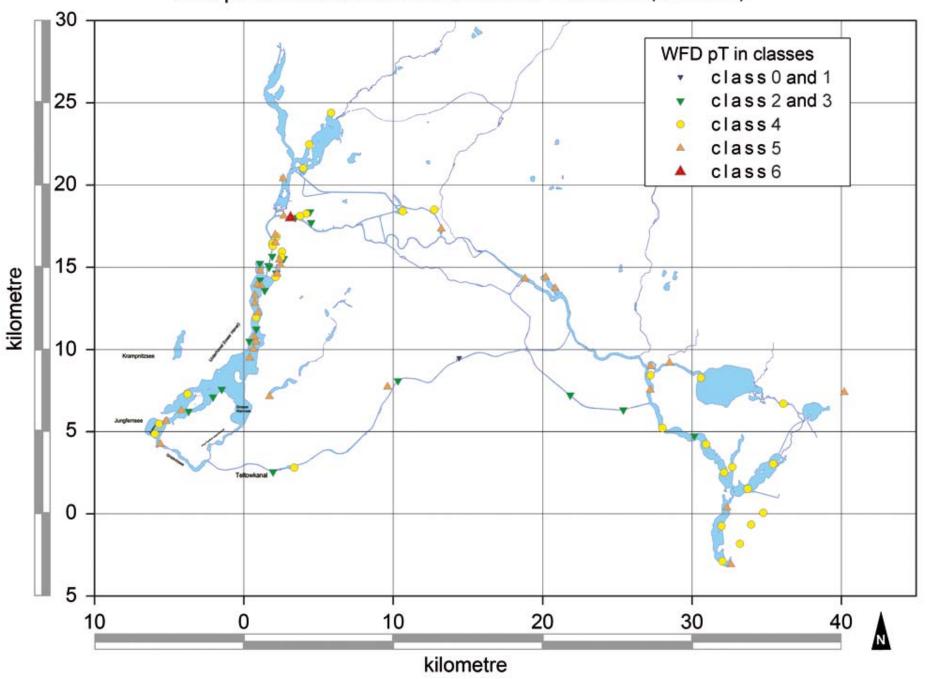
Toxicity Classification / LID (Lowest identified dilution step with no effect)

Elutriates of River Sediments:17β-Estradiol Equivalents [µg/l]



Highest			Toxicity classes		Management	categories
dilution level without effect	Dilution factor	pT-value	7-level system	Designation	3-level assessment	Colour coding
Original sample	20	0	0	toxicity non-detectable		0
1:2	2 <sup>-1</sup>	1	Ι	very slightly toxic	uncritically polluted	T
1:4	2-2	2	II	slightly toxic		П
1:8	2 <sup>-3</sup>	3	III	moderately toxic	critically	Ш
1:16	2-4	4	IV	distinctly toxic	polluted	IV
1:32	<b>2</b> -5	5	V	highly toxic	hazardana	V
≤ (1:64)	≤ <b>2</b> -6	≥ 6	VI	extremely toxic	hazardous	VI

WFD: pT-values and Classification of River Sediments (Elutriates)



Environmental Monitoring and Assessment (2005) DOI: 10.1007/s10661-005-9061-0

© Springer 2005

#### ECOTOXICOLOGICAL CLASSIFICATION OF THE BERLIN RIVER SYSTEM USING BIOASSAYS IN RESPECT TO THE EUROPEAN WATER FRAMEWORK DIRECTIVE

GERD HUSCHEK and P.-D. HANSEN\*

Department of Ecotoxicology, Technische Universität Berlin, Faculty VI, Franklin Strasse 29 (OE4), D-10587 Berlin, Germany

(\*author for correspondence, e-mail: pd.hansen@tu-berlin.de)

(Received 21 February 2005; accepted 7 October 2005)

Abstract. Bioassays as well as biochemical responses (biomarkers) in ecosystems due to environmental stress provide us with signals (environmentally signalling) of potential damage in the environment. If these responses are perceived in this early stage in ecosystems, the eventual damage can be prevented. Once ecosystem damage has occurred, the remedial action processes for recovery could be expensive and pose certain logistical problems. Ideally, "early warning signals" in ecosystems using sensing systems of biochemical responses (biomarkers) would not only tell us the initial levels of damage, but these signals will also provide us with answers by the development of control strategies and precautionary measures in respect to the European Water Framework Directive (WFD). Clear technical guidelines or technical specifications on monitoring are necessary to establish and characterise reference conditions for use in an ecological status classification system for surface water bodies. For the Ecotoxicological Risk Assessment (ERA) of endocrine effects we used an approach of the exposure - dose - response concept. Based on the "Ecototoxicological Classification System of Sediments" that uses pT-values to classify effects in different river systems, we transferred the biomonitoring data to the five-level ecological system of the WFD. To understand the complexity of the structure of populations and processes behind the health of populations, communities and ecosystems an ERA should establish links between natural factors, chemicals, and biological responses so as to assess causality. So, our ecological monitoring assessment has incorporated exposure & effects data.

Keywords: bioassay, effect assessment, ecotoxicological classification in sediment, endocrine effects, exposure of drugs

#### 1. Introduction

For risk assessment and risk assessment tools new recommendations are described in the Technical Guidance Document of EU – Edition 2 (TGD, 2003), in the new EU Chemicals Legislation REACH (REACH, 2004) and in the status report for toxicological methods of the European Centre for the Validation of Alternative Methods (ECVAM, 2003). In the EU Water Framework Directive (WFD, 2000) a general requirement for ecological protection, and a general minimum chemical standard, was introduced to cover all surface waters. For the description of "good ecological status" and "good chemical status" effects monitoring tools are needed for the description of the ecological status of river basin systems. In some case

# Exposure Toxicity Ratio = ETR

# **Exposure-Toxicity-Ratio:** *ETR*

The **risk potential** will be expressed as a ratio of the **exposure** (Predicted Environmental Concentration: *PEC*) and the **toxicity of the active substance** (*LC50*).

acute risk potential

chronic risk potential

$$ETR_{acute} = \frac{sPEC}{LC50_{species}}$$

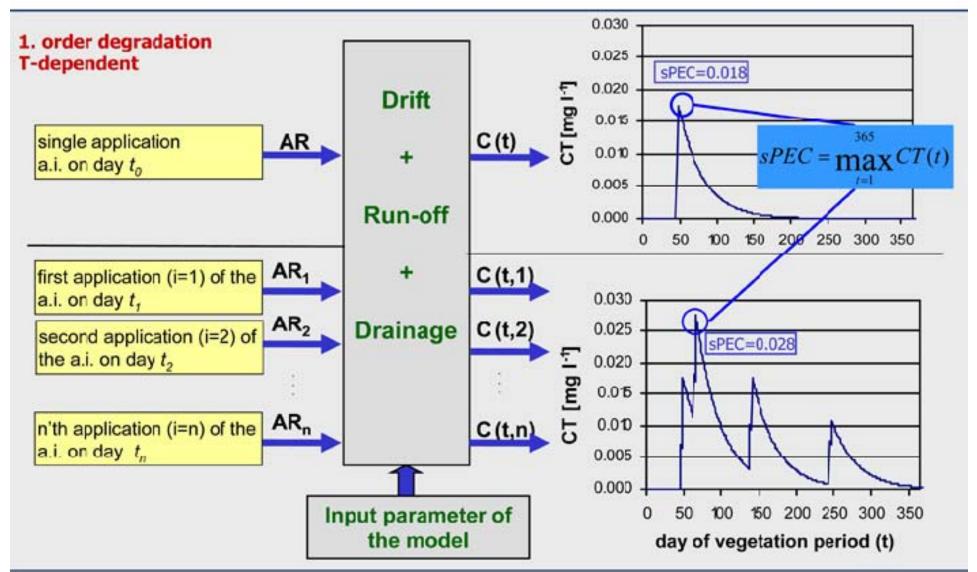
$$ETR_{chronic} = \frac{lPEC}{NOEC_{species}}$$

ETR <sub>species</sub>	Exposure-Toxicity-Ratio
sPEC	short-term exposure in the surface water
IPEC	long-term exposure in the surface water
LC50 <sub>species</sub>	Lethal concentration
NOEC species	No effect concentration





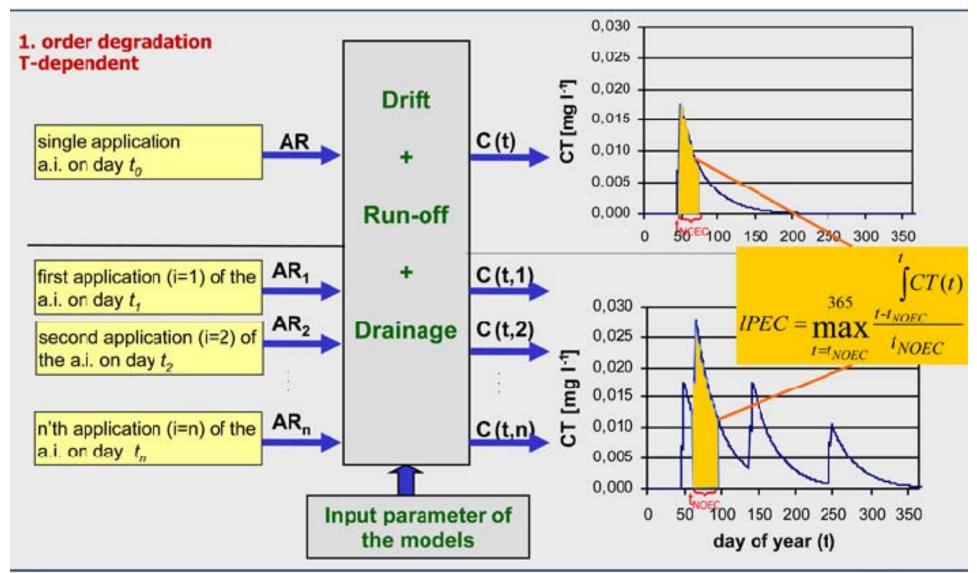
### Calculation of *sPEC*: acute risk







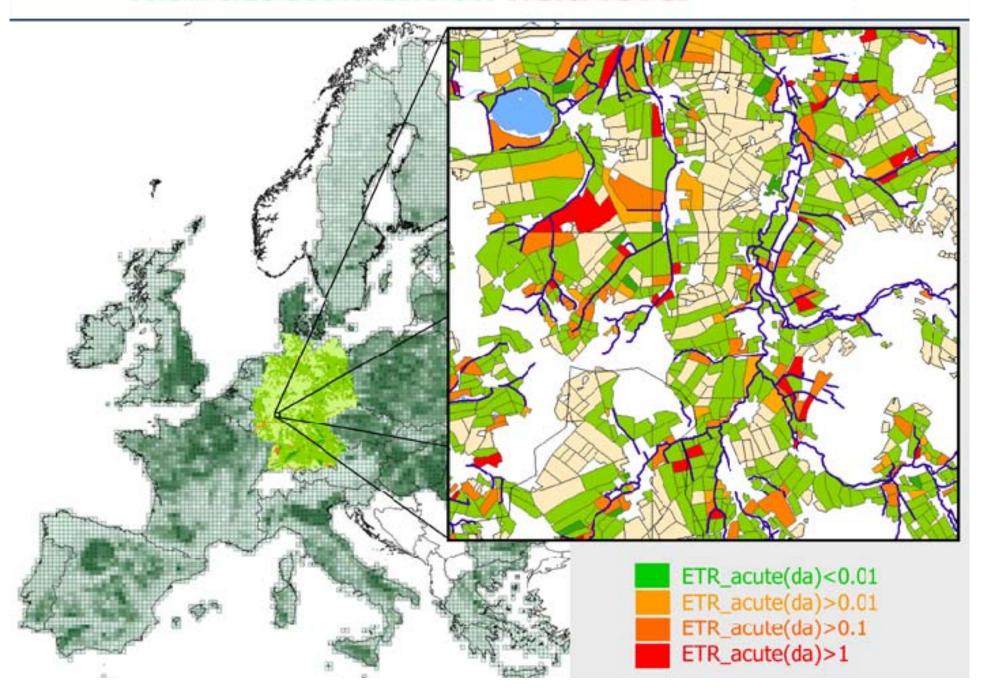
## Calculation of *IPEC*: chronic risk







# Risk assessment on field level

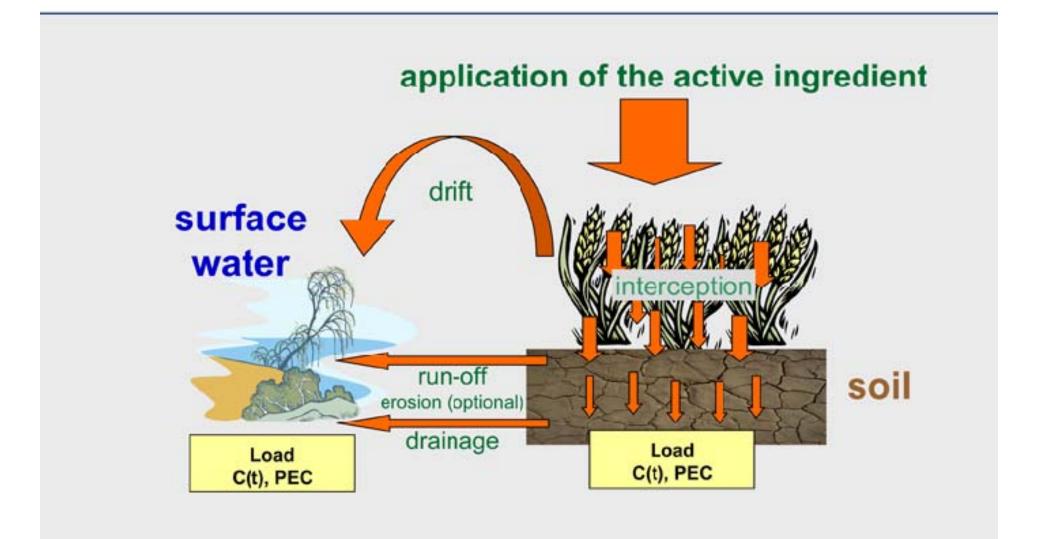


# Spatial aggregation of acute risk (*ETR*<sub>acute(daphnia)</sub>)

Risk potentials *ETR*<sub>acute(daphnia)</sub> were calculated without considering buffer zone distances.

fraction of arable land weighted mean fraction of arable land with ETR<sub>acute</sub>>0.1 (by area) with ETR<sub>acute</sub>>1 0.0000 - 0.0010

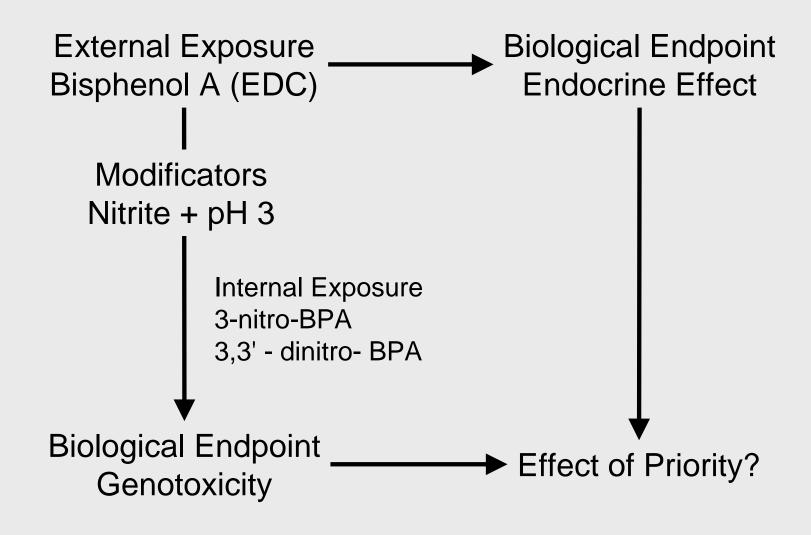
# **Modeled pathways**

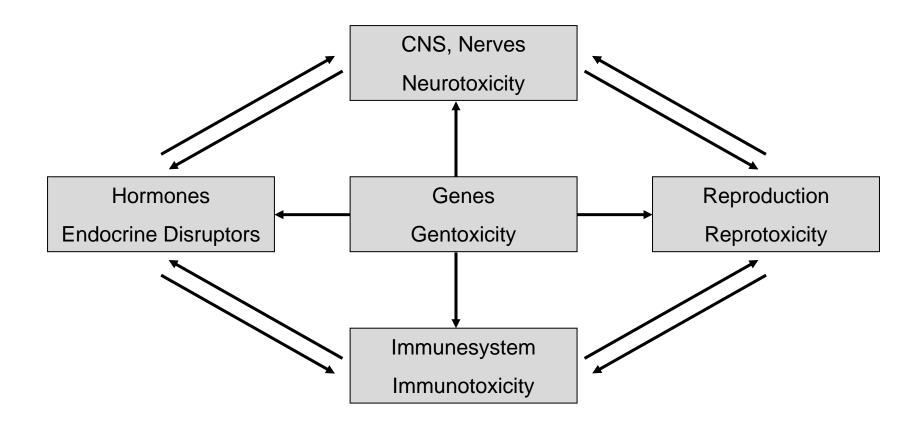






Realistic Exposure Szenarios and Modificators (mixed effects)





**Quality Norms: QN, UQN, EQN** 

Single Substance Approach



# Tools = Bioassays

Approach used to derive Quality Norms (UQN/EQN) for aquatic communities (Recognized test methods include methods developed by DIN, ISO, CEN, OECD)

**General principle**: lowest test result for the most sensitive species is to be used as the starting point for the derivation of quality norms (QN).

#### Compensationfactors

(uncertainty - extrapolation test results to the real environment)

- F1 = 0.1 NOEC data are available for each trophic stages
  F2 = 0.01 NOEC data are available for at least two of the trophic stages
  in addition, a wide range of aquatic toxicity data is available
- F2 = 0.1 additional risk:
   the substance is persistent in the aquatic environment (BCF > 100)
   the substance is capable of transforming into dagerous substances synergism and/or additive effects

```
Quality Norm (QN): Fishery

QO [μg/L] = WF [mg/kg]
------
BCF [l/kg]
```

Quality Norm (QN): supply of drinking water

QO = 0.5 TG (TG = Limit value for drinking water)

(For the xenobiotic substances listed in Annex 2 of the Drinking Water Ordinance, the reduction factor F may not exeed

#### **Tools WFD Environmental Quality Norm (EQN)**

**Standardisation and Normalisation DIN, CEN, ISO 2006:** 

Effect related parameters already in the regulations: Bacteria-, Algae-, Crustaceans-, Fish- Assays, Genotoxicity Replacements: Fish Egg Assay, DIN 38415 T6 / ISO NWI

Genotoxicity (procaryontic assays):
umu-Assay, DIN 38415 T3 - ISO 13829 (Gov.Lab.)
AMES-Assay, DIN 38415 T4 - ISO/CD 16240 (Industry)
In addition:
Eucaryontic assays (except COMET assays):
Micronulei-Assay, ISO/CD 21427

New effect related parameters for regulations:

- (1) Immunotoxicity, DIN UA 5.7
- (2) Endocrine effects, DIN UA 7.3 / ISO TC147/SC5 NW

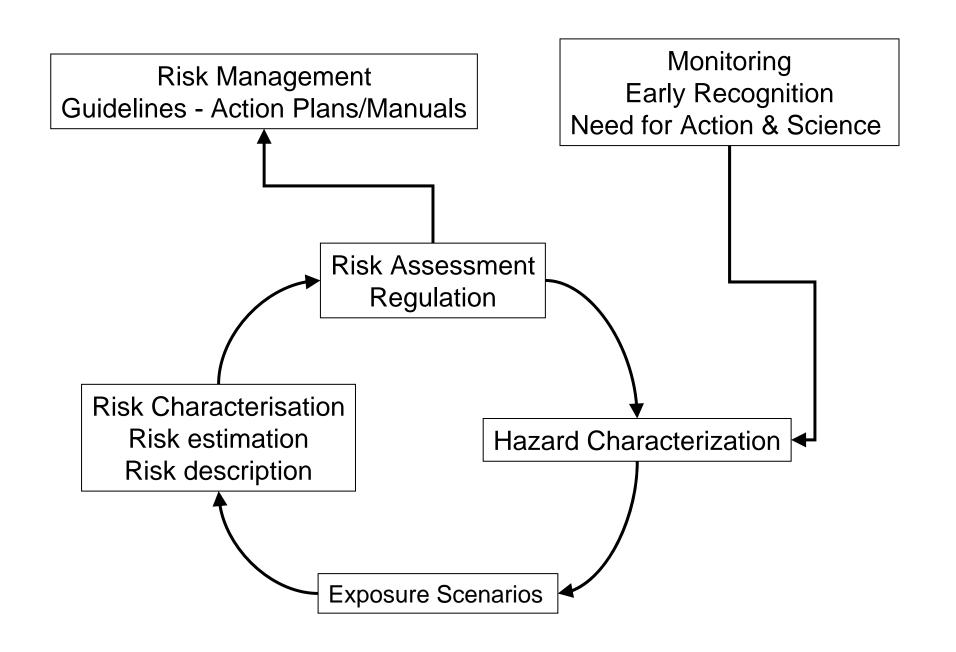
# Quality Norm (QN) for PCBs of the Lower Havel

EG-Nr.	Stoff	QN	Mittel 1999	Mittel 2000	Mittel 2001
99e	Fluoranthen	0,025 µg/l	<qz< td=""><td>0,028</td><td>0,027</td></qz<>	0,028	0,027
101a	PCB 101	20 μg/kg	n.g.	45,27	32,76
101b	PCB 118	20 μg/kg	n.g.	n.g.	27,30
101c	PCB 138	20 μg/kg	n.g.	55,96	39,82
101d	PCB 153	20 μg/kg	n.g.	52,52	31,80
101e	PCB 180	20 μg/kg	n.g.	34,04	22,93
101f	PCB 28	20 µg/kg	n.g.	91,76	32,76
101g	PCB 52	20 μg/kg	n.g.	40,50	30,35

Compound	76/464/EWG <b>a) List I</b>	QN	Rili 76/464	QN	QN
	b) List II			Water	Suspended Solids
ННСВ	BW, BE, BY, TH			7,0 μg/l	
loxynil	List II		0,1 μg/l	0,1 μg/l	
Kobalt	List II		80 mg/kg	0,9 μg/l	50 mg/kg
Molybdän	List II		5 mg/kg	7,0 μg/l	8,9 mg/kg
Monolinuron	List I – 99S	0,1 μg/l		0,01 μg/l	
Propanil	List I – 99S	0,1 μg/l		0,9 μg/l	
Propazin	List II		0,1 μg/l	0,25 μg/l	

Compound	QN in μg/l [g/kg
Acenaphthen	0,32[0,08]
	0,32
	[0,08]
Anilin	0,81
Benz(a)anthracen	0,002
	0,002
Bezafibrat	
Butylbenzylphthalat	5,2
Carbamazepin	0,5
Chlorbenzilat	0,6
Chrysen	0.9
Clofibrinsäure	5
Cyanazin	0,12
Desethylterbutylazin	
Desisopropylatrazin	
Desmetryn	0,03
Dibutylphthalat	10
Dibutylzinn	0,21

Diclofenac	0,1
EDTA-Na	2200
Ethofumesat	24
Fluor	
Fluoren	2,1[0,9]
	2,1
	[0,9]
Fluroxypyr	152
lopamidol	
Kresoxim-methyl	0,3
Lenacil	1
Metalaxyl	120
Methyl-tertbutylether (MTBE)	2600
Methylisothiocyanat	0,05
Metobromuron	2
Metoxuron	0,09
Metribuzin	0,18
NTA / NTA-Natriumsalz	930
2-Methoxyanilin (o-Anisidin)	5,5
Pencycuron	1,3[1,3]
Pendimethalin	0,27
Penanthren	0.5
	+



# Which substances are we talking about?

- 76/464/EEC
  - Dangerous substances
  - "List I"
    - The 18 substances regulated under "so called daughter Directives"
    - "PBT"
  - "List II"
    - All other substances belonging to the categories included in Annex I (List I and List II)
    - Deleterious effect on the aquatic environment
    - Metals
    - Identified by the Member States

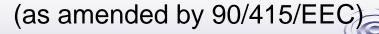
- Water Framework Directive (2000/60/EC)
  - Priority substances:
    - "presenting a significant risk to or via the aquatic environment"
    - included in annex X WFD
  - Hazardous substances (or priority hazardous substances) :
    - PBT or other equivalent concern
    - Part of the priority substances in annex X WFD
  - Other pollutants :
    - "any substance liable to cause pollution"
    - Identified by river basin
    - categories included in annex VIII

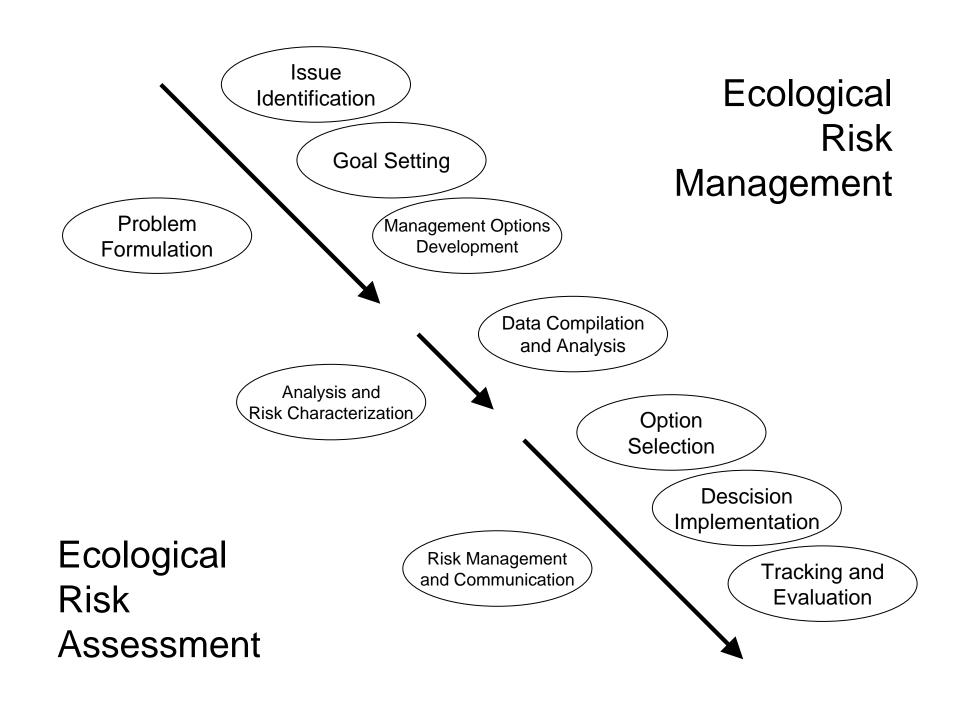
# List of 33 Priority Substances (Decision 2455/2001/EC or Annex X WFD)

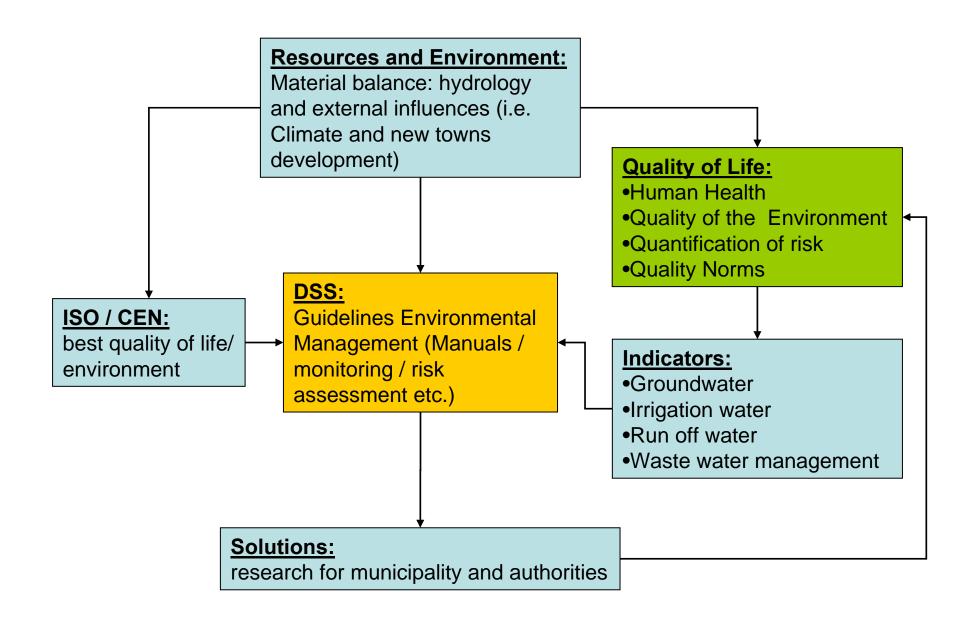
PHS under review **Priority Priority hazardous** substances substances Anthracen Atrazine • Cadmium Chlorpyrifos Alachlor • C10-13-chloroalkanes DEHP • Benzene • Hexachlorobenzene • Diuron • Chlofenvinphos Hexachlorobutadiene • Endosulfan • 1,2-Dichloroethane Hexachlorocyclohexane • Isoproturon • Dichloromethane Mercury • Lead Flouranthene Nonylphenols Naphtalene Nickel and its PAH • Octylphenols compounds Pentachlorobenzene Pentachlorophenols Trichloromethane • PBDE • Simazine (Chloroform) Tributyltin compounds • Trichlorobenzenes Trifluralin

## "Daughter Directives"

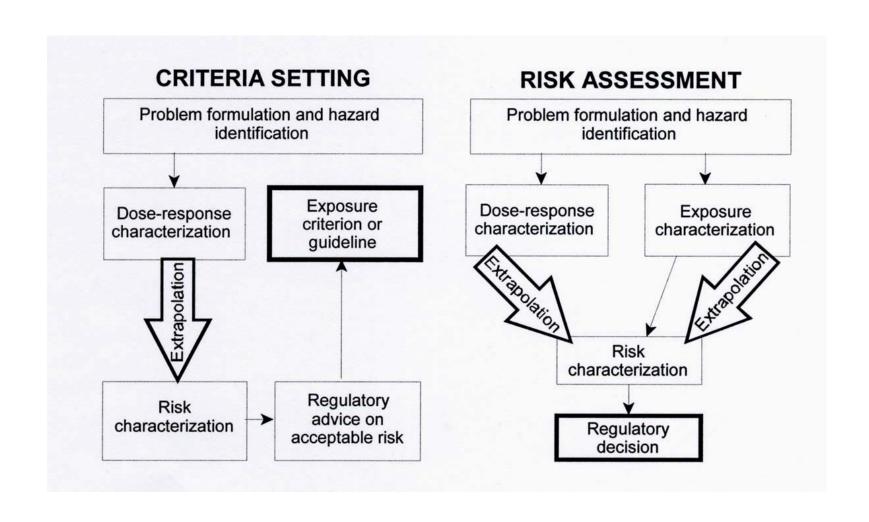
- 82/176/EEC: Mercury (by chlor-alkali electrolysis industry)
- 83/513/EEC : Cadmium
- 84/156/EEC : Mercury (by other sectors)
- 84/491/EEC : Hexachlorocyclohexane
- 86/280/EEC : Dangerous Substances Discharges :
  - Carbon tetrachloride
  - DDT
  - Pentachlorophenol
  - Aldrin, Dieldrin, Endrin and Isodrin (as amended by 88/347/EEC)
  - Hexachlorobenzene
  - Hexachlorobutadiene
  - Chloroform
  - 1,2-dichloroethane (EDC)
  - Trichloroethylene (TRI)
  - Perchloroethylene (PER)
  - Trichlorobenzenes and 1,2,4-TCB



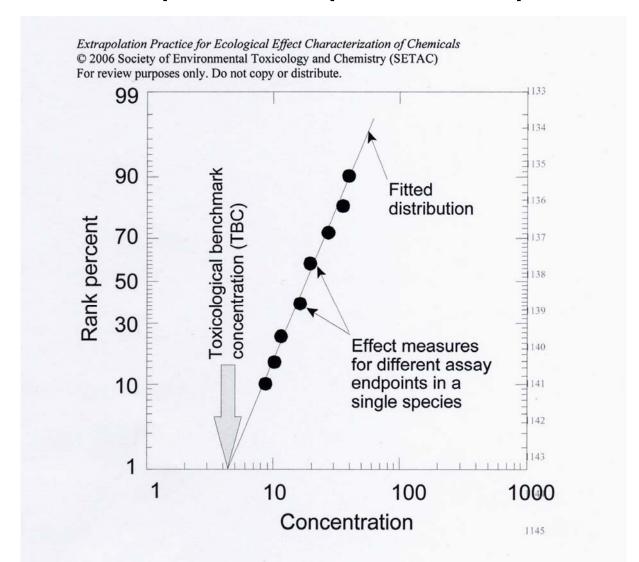




### Setting criteria and assessing risk from existing exposures



### Distribution of endpoints for a species to extrapolate a TBC



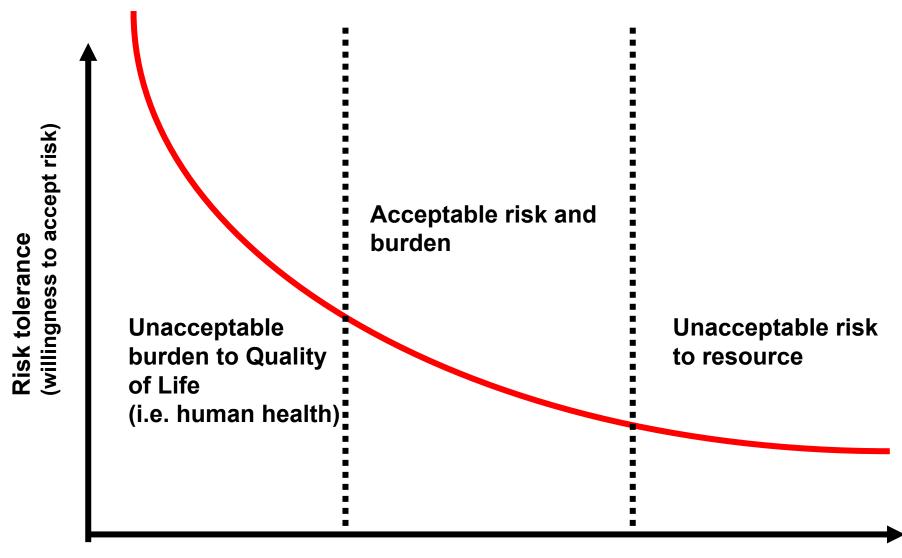
Toxicological benchmark concentration (TBC)

Benchmarking may be defined as "a continuous process to look for the best operating practices in order to achieve the most superior (environmental) performance".

Other definitions have been given in the literature, but this is the one closest to the workfield of Environmental Investigations.

There are three main items in this definition:

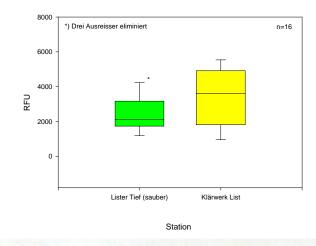
- 1. Environmental performance
- 2. The best operating practices
- 3. a continuous process
- toxicity load as such;
- toxicity load per tons of production capacity;
- toxicity load per tons of product really produced



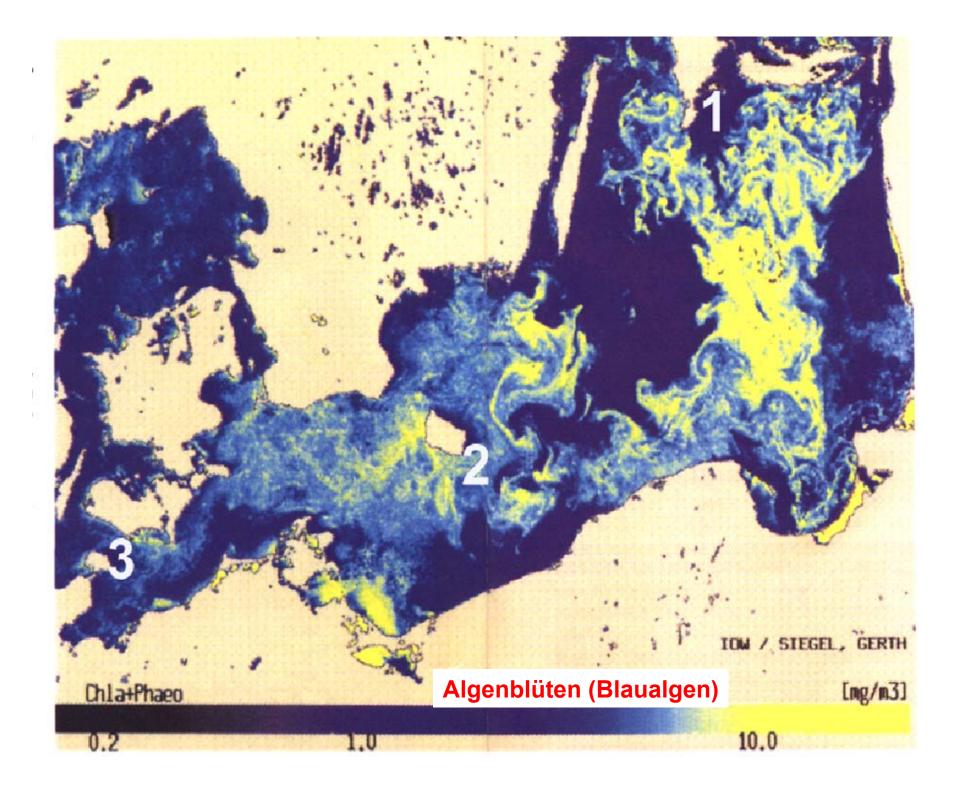
Perceived value of an ecological Resource



### Phagozytosis activity of exposed *M. edulis* Island of Sylt, May 2005







## Overview of Key Phytoplankton Toxins and Their Recent Occurrence in the North and Baltic Seas

B. Luckas, J. Dahlmann, K. Erler, G. Gerdts, N. Wasmund, C. Hummert, P. D. Hansen<sup>5</sup>

Received 3 March 2004; revised 4 August 2004; accepted 1 September 2004

ABSTRACT: The frequency and intensity of harmful algal blooms (HABs) appear to be on the rise globally. There is also evidence of the geographic spreading of toxic strains of these algae. Consequently, methods had to be established and new ones are still needed for the evaluation of possible hazards caused by increased algal toxin production in the marine food chain. Different clinical effects of algae-related poisoning have attracted scientific attention; paralytic shellfish poisoning, diarrhetic shellfish poisoning, and amnesic shellfish poisoning are among the most common. Additionally, cyanobacteria (blue-green algae) in brackish waters often produce neurotoxic and hepatotoxic substances. Bioassays with mice or rats are common methods to determine algal and cyanobacterial toxins. However, biological tests are not really satisfactory because of their low sensitivity. In addition, there is growing public opposition to animal testing. Therefore, there has been increasing effort to determine algal toxins by chemical methods. Plankton samples from different European marine and brackish waters were taken during research cruises and analyzed on board directly. The ship routes covered marine areas in the northwest Atlantic, Orkney Islands, east coast of Scotland, and the North and Baltic seas. The first results on the occurrence and frequency of harmful algal species were obtained in 1997 and 1998. During the 2000 cruise an HPLC/MS coupling was established on board, and algal toxins were measured directly after extraction of the plankton samples. In contrast to earlier cruises, the sampling areas were changed in 2000 to focusing on coastal zones. The occurrence of toxic algae in these areas was compared to toxin formation during HABs in the open sea. It was found that the toxicity of the algal blooms depended on the prevailing local conditions. This observation was also confirmed by monitoring cyanobacterial blooms in the Baltic Sea. Optimal weather conditions, for example, during the summers of 1997 and 2003, favored blooms of cyanobacteria in all regions of the Baltic. The dominant species regarding the HABs in the Baltic was Nodularia spumigena. However, in addition to high concentrations of Nodularia spumigena in coastal zones, other blue-green algae are involved in bloom formation, with changes in plankton communities influencing both toxin profiles and toxicity. © 2005 Wiley Periodicals, Inc. Environ Toxicol 20: 1–17, 2005.

Keywords: algal toxins; paralytic shellfish poisoning (PSP) toxins; domoic acid; diarrhetic shellfish poisoning (DSP) toxins; spirolides; nodularin; LC/MS analysis

#### HARMFUL ALGAL BLOOMS

Correspondence to: P. D. Hansen; pd.hansen@tu-berlin.de
Published online in Wiley InterScience (www.interscience.wiley.com).
DOI 10.1002/tox.20072

Natural "contaminants" may accumulate via food chains. A typical example is the ingestion and accumulation of algal toxins in filter feeders such as mollusks. Because mollusks

@ 2005 Wiley Periodicals, Inc..

<sup>&</sup>lt;sup>1</sup>Institute of Nutrition, University of Jena, Jena, Germany

<sup>&</sup>lt;sup>2</sup>Biologische Anstalt Hellgoland, Alfred Wegener Institute for Polar and Marine Research, Helgoland, Germany

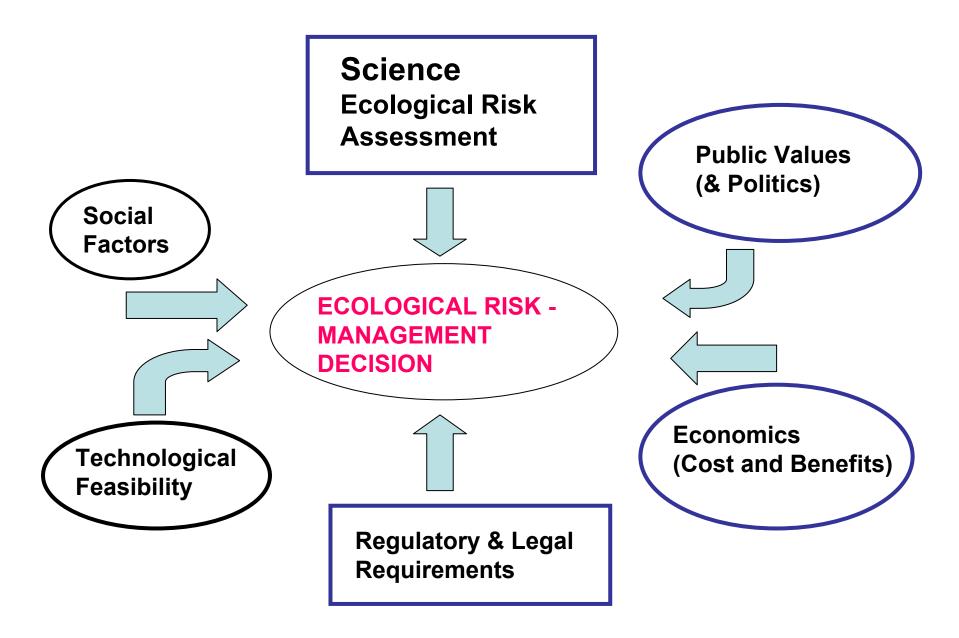
<sup>3</sup>Institute of Baltic Research, Warnemuende, Germany

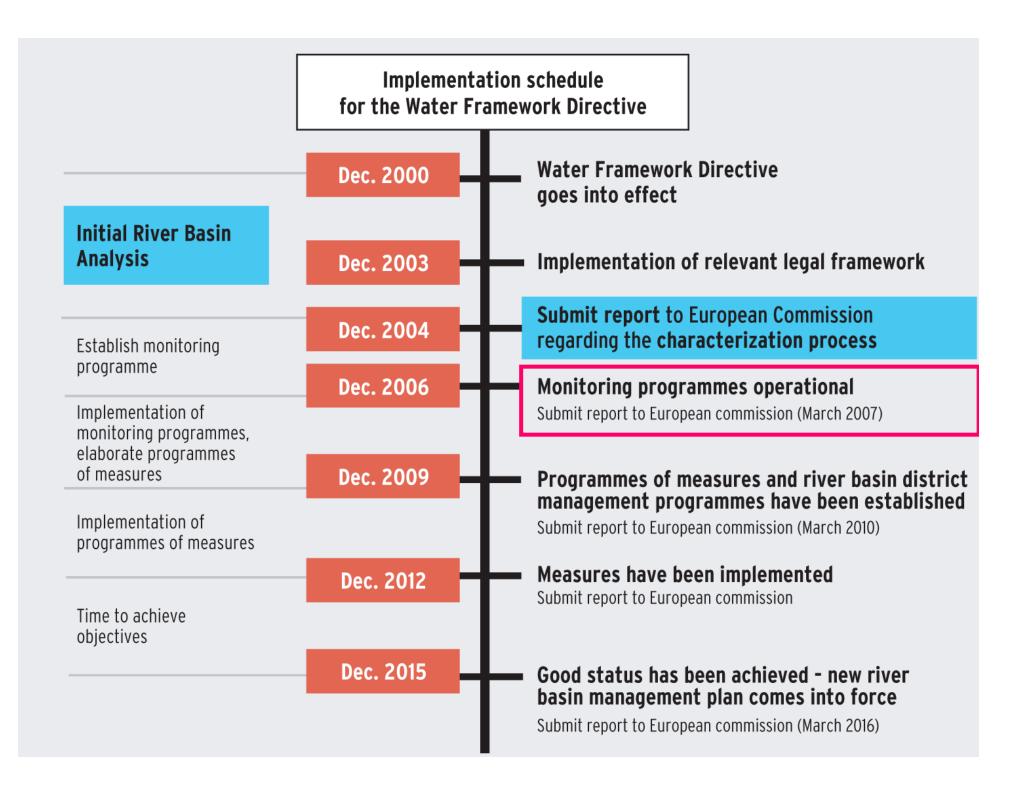
<sup>&</sup>lt;sup>4</sup>Labor WEJ (Eurofins), Hamburg, Germany

<sup>&</sup>lt;sup>5</sup>Institute for Ecological Research and Technology, Department of Ecotoxicology, Technische Universitaet Berlin (Berlin University of Technology), Berlin, Germany



### Inputs to the ecological risk management decisions





### The WFD describes:

### Surveillance monitoring

The assessment and description of long-term ecological trends and an overall description of the waters to determine whether a good status has been or will be achieved.

### Operational monitoring

The assessment of the status of the water mass of which it has become evident that it may not meet the environmental objectives and/or to assess the changes in the status of this water mass resulting from the programme of measures.

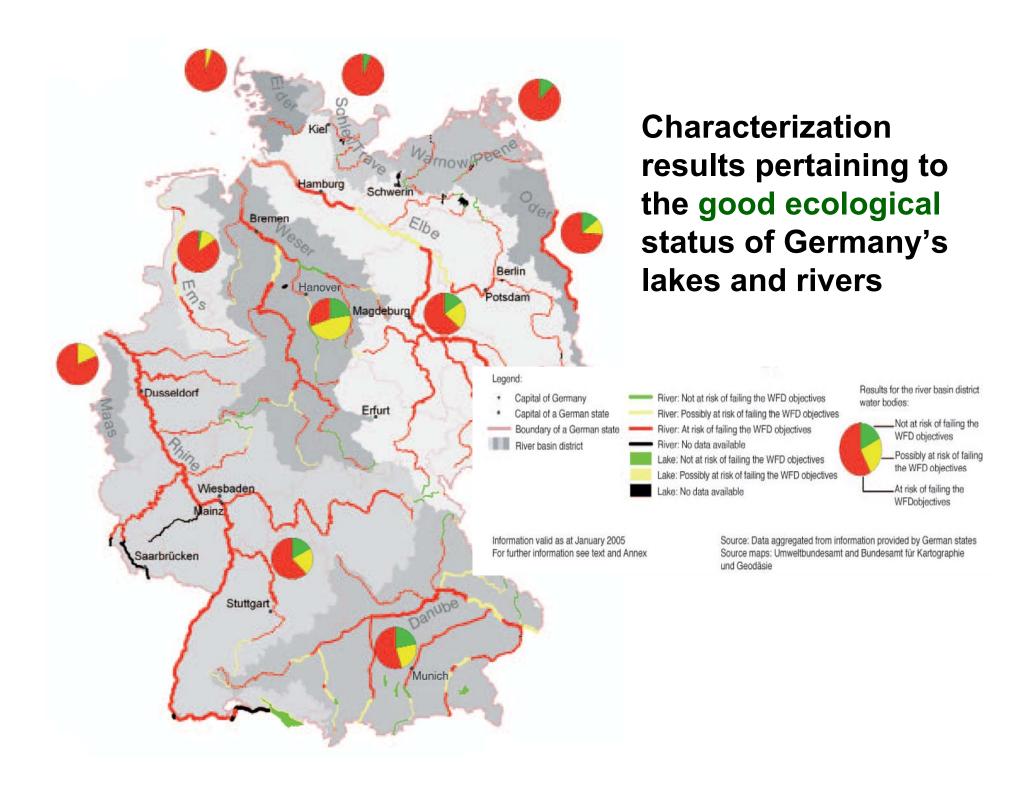
### Investigative monitoring

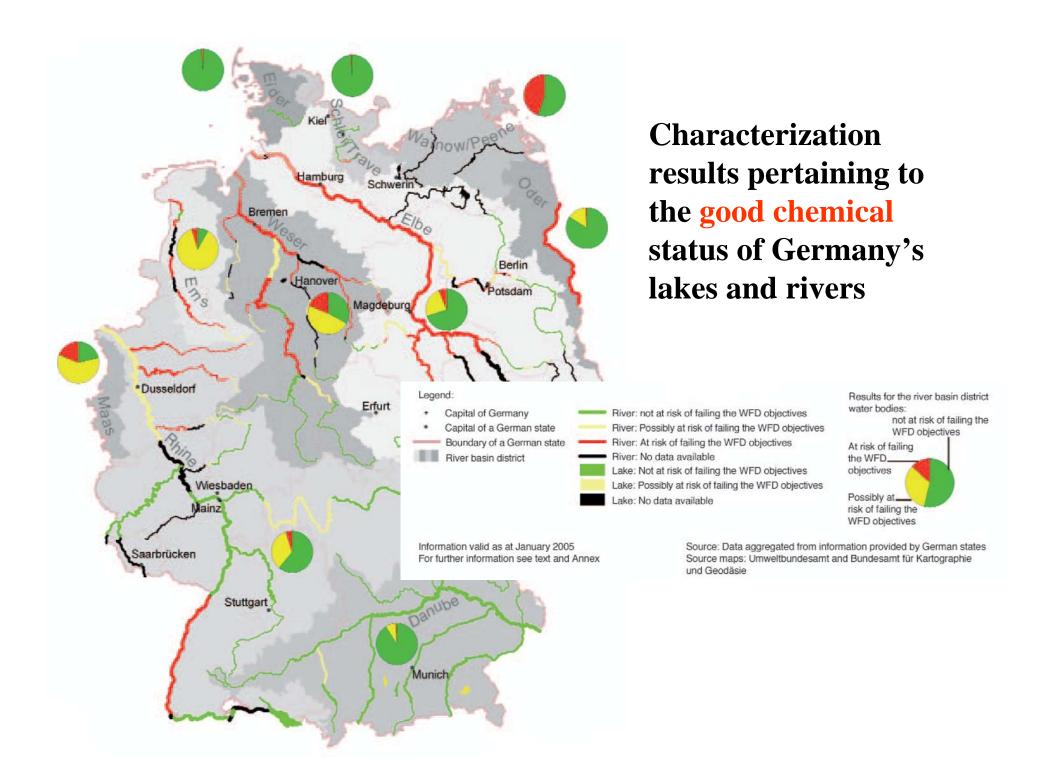
The identification of the cause of the failure to achieve a good status.

### The characterization process and the three questions:

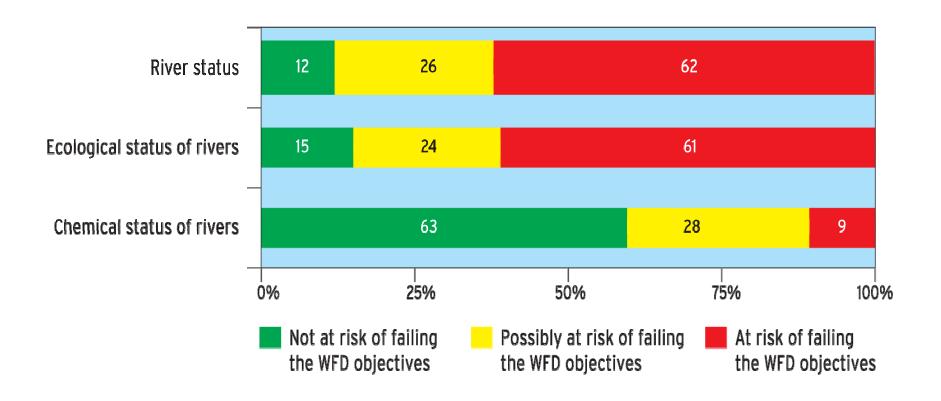
- 1. Which water bodies might fail the environmental WFD objectives
- 2. Which chemical and non-chemical pressures are to blame for the failure to meet these objectives?
- 3. Which mechanisms and effects should therefore be the focus of operative monitoring

The results of these investigations will form the basis for the elaboration of the monitoring programmes

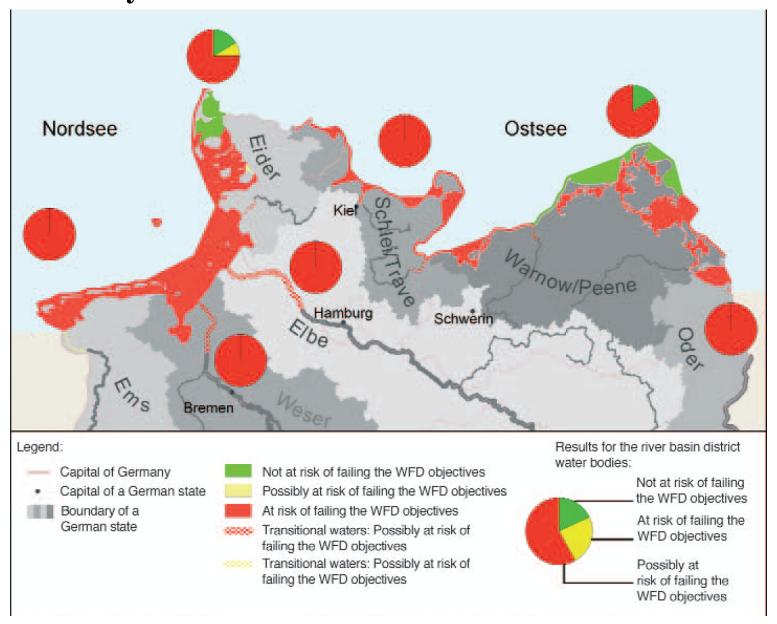




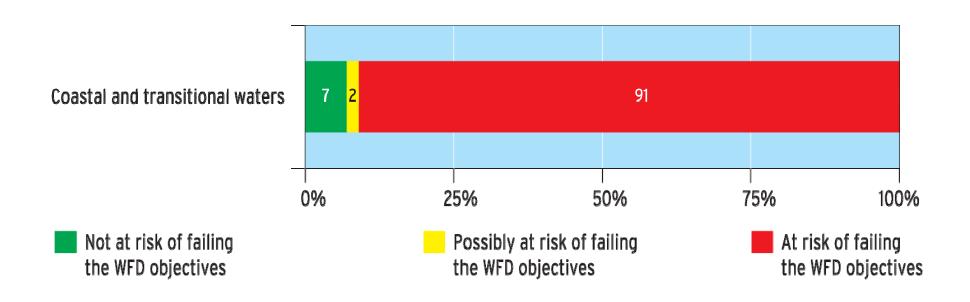
### Characterization results for rivers



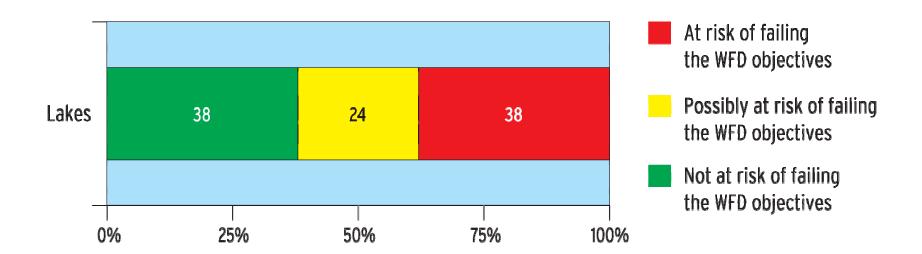
# Characterization results pertaining to the good status of Germany's transitional and coastal waters



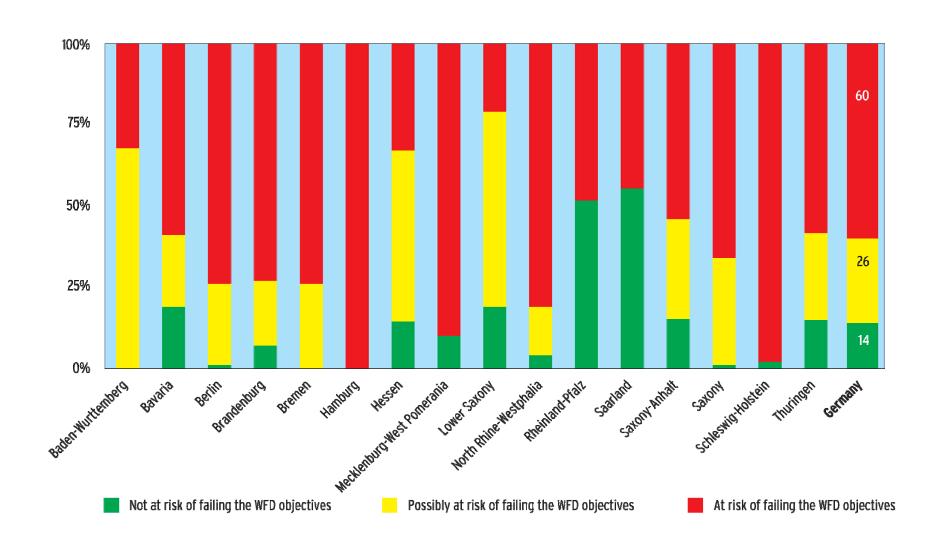
# Characterization results for coastal and transitional waters



### **Assessment results for lakes**



### Initial characterization of Germany's surface water bodies



### Environmental Quality Standards (EQS) for Priority Substances in Suface Water (AA: Annual Average; MAC: Maximum Allowable Concentration)

Unit: [µg/l].

(1)	(2)	(3)	(4)	(5)	(6)	(7)
N°	Name of substance	CAS number	AA-EQS Inland surface waters	AA-EQS Other surface waters	MAC- EQS Inland surface waters	MAC-EQS Other surface waters
(1)	Alachlor	15972-60-8	0.3	0.3	0.7	0.7
(2)	Anthracene	120-12-7	0.1	0.01	0.4	0.4
(3)	Atrazine	1912-24-9	0.6	0.6	2.0	2.0
(4)	Benzene	71-43-2	10	8	50	50
(5)	Pentabromodiphenylether	32534-81-9	0.0005	0.0002	1.4	1.4

## Environmental Quality Standards (EQS) for Priority Substances in Surface Water (AA: Annual Average; MAC: Maximum Allowable Concentration)

Unit:  $[\mu g/l]$ .

(1)	(2)	(3)	(4)	(5)	(6)	(7)
N°	Name of substance	CAS number	AA-EQS Inland surface waters	AA-EQS Other surface waters	MAC- EQS  Inland surface waters	MAC-EQS Other surface waters
(1)	DDT total	50-29-3	0,025	0,025	not applicable	not applicable
	para-para-DDT	CHECK	0,01	0,01	not applicable	not applicable
(2)	Aldrin	309-00-2	S=0,010	S=0,005	not applicable	not applicable
(3)	Dieldrin	60-57-1				
(4)	Endrin	72-20-8				
(5)	Isodrin	465-73-6				
(6)	Carbontetrachloride	56-23-5	12	12	not applicable	not applicable
(7)	Tetrachloroethylene	127-18-4	10	10	not applicable	not applicable
(8)	Trichloroethylene	79-01-6	10	10	not applicable	not applicable

#### Änderungen AA-EQS für Fließgewässer

Nr	Stoff	AA-EQS Juli 05	AA-EQS Dez. 05	AA-EQS Mai 06	ZV A
16	НСВ	0,0004	0,0002	0,01	0,01
17	Hexachlorbutadien	0,003	0,003	0,1	0,5
20	Blei	2,1	2,1	7,2	3,4 1)
23	Nickel	1,7	3,8	20	4,4 1)
25	Octylphenol	0,06	0,06	0,1	
27	Pentachlorphenol	0,2	0,4	0,4	bereits Dez.05 geändert
28	Summe Benzo(g,h,i)perylen Indeno(1,2,3-cd)pyren	0,02	0,002 2)	0,002 2)	bereits Dez.05 geändert
32	Trichlormethan	12	2,5	2,5	bereits Dez.05 geändert

#### Änderungen MAC-EQS für Fließgewässer

Nr	Stoff	MAC-EQS Juli 05	MAC-EQS Dez. 05	MAC-EQS Mai 06	
16	НСВ	0,05	0,002	0,05	
17	Hexachlorbutadien	0,6	0,04	0,6	
19	Isoproturon	1,3	1	1	bereits Dez.05 geändert
28	Benzo-a-pyren	0,05	0,1	0,1	bereits Dez.05 geändert
30	ТВТ	0,002	0,0015	0,0015	bereits Dez.05 geändert

green = AAEQS yellow <10%; orange=10-25%; red >25%

### Summary

The good ecological status is mainly supported by the hydromorphological elements (no Bioassays!)

Comprehensive strategy is given for a Ecological risk assessment (ERA) that based on two major elements: characterization of effects and characterization of exposure.

Effect related appproaches (bioassays, biomarkers, biosensors (?) - environmental signalling (on line real time bioassays) - are only relevant in the context to the "Quality Norms (QN)" developed after the WFD in correspondence to the "good chemical status"

It was demonstrated how to develop a strategy of classification of sediments and how to include effect related data of river sediments into the WFD concept.

Measurement data from monitoring programmes provide the basis for water quality evaluations (QN) for the purpose of achieving the environmental objectives of the Water Framework Directives (WFD).

**Further tools (Swift Programm):** 

Contribution of the REACH Concept for Natural Water Conservation?!

REACH=Registration, Evaluation, Authorisation of Chemicals (Registration, Bewertung, Zulassung und Beschränkung von chemischen Stoffen)

**QSAR=Qualitative Structure Activity Relationship** 

If there is a river "at risk"?

Monitoring in support of ERA should be adequate for the characterization of exposure and effects thus enabling the sustainable development for aquatic life and human health protection.

### **Acknowledgement**

**European Commission, DG Research I.-3** 

**SedNet, European Sediment Research Network** 



CITY FISH: Modelling the Ecological Quality of Urban Rivers: Ecotoxicological Factors Limiting Restoration of Fish Populations

SANDRINE - www.sandrine-wwc.de

**BEEP, BIOMAR I+II** 





## **Thank You!**

www.tu-berlin.de/~oekotox www.ecosystemhealth.net



### Literature:

Hansen, P.-D. 2002. Unerwünschte Wirkungen. In: A. Grohmann (Hrsg.), 8. Auflage Karl Höll: Wasser, Nutzung im Kreislauf, Hygiene, Analyse und Bewertung, Walter de Gruyter, Berlin, New York, 550-563

Hansen, P.-D. 2003. Biomarkers. In: B.A. Markert, A.M. Breure and H.G. Zechmeister (Eds), Bioindicators & Biomonitors, Principles, Conceps and Applications, Elsevier, Amsterdam, Boston, London, Oxford, Paris, San Diego, San Francisco, Singapore, Sydney, Tokyo, 203-220

Garcia-Reyero, N., Requena, V, Petrovic, M., Fischer, B., Hansen, P.-D., Diaz, A. Ventura, F., Barcelo, D. and Pina, B. 2004 Estrogenic Potential of Halogenated Derivates of Nonylphenol Ethoxylates and Carboxylates. Environmental Toxicology and Chemistry, 23, 3, 705-711

Huschek, G., Hansen, P.-D., Maurer, H., Krengel, D., Kayser, A. 2004 Environmental Risk Assessment of medicinal products for human use in respect to the recommendations of the European Commission. Environmental Toxicology, Vol. 19, 3, 226-240 Luckas, B., J. Dahlmann, K.Erler, G. Gerdts, N. Wasmund, C. Hummert, P.-D. Hansen. 2005. An Overview on Key Phytoplankton Toxins and their recent occurence in North Sea and Baltic. Environmental Toxicology, 20,1, 1-17

Huschek, G. and P.-D. Hansen. 2006. Ecotoxicological classification of the Berlin river system using bioassays in respect to the European Water Framework Directive. Environmental Monitoring and Assessment, 121, 1-3

Gagné, F., C. Blaise, M. Fournier, P.D. Hansen. 2006. Effects of selected pharmaceutical products on phagocytic activity in Elliptio complanata mussels. Comp. Biochem. Physiol. C Toxicol Pharmacol, 143 (2), 179-186

Arab, N, S. Lemaire-Gony, E. Unruh, P.D. Hansen, B.K. Larsen, O.K. Andersen, N.J. Narbonne 2006.

Preliminary study of responses in mussels (Mytilus edulis) exposed to bisphenol A, diallyl phthalate and tertebromodiphenyl ether. Aquat Toxicol. 78 Suppl 1, 86-92

P.-D. Hansen, J. Blasco, A. De Valls, V. Poulsen, M. van den Heuvel-Greve, 2006. Biological analysis (Bioassays, Biomarkers, Biosensors) In: Sustainable management of sediment resources, Volume 2, Sediment quality and impact assessment of pollutants. Eds. Damia Barceló and Mira Petrovic, Elsevier Publishers Amsterdam, London, New York, 311 pp (in press)

EG-Wasserrahmenrichtlinie: Interaktiver Bericht zur Bestandsaufnahme in Deutschland, DVD Auflage Februar 2006, Umweltbundesamt

Wasserrahmenrichtlinie: Dokumente zur EG-Wasserrahmenrichtlinie, DVD Auflage Februar 2006, Umweltbundesamt

Blondzik, K. et al. Wasserwirtschaft in Deutschland Teil 2 Gewässergüte. Hrsg. BMU: Umweltpolitik, Wasserwirtschaft in Deutschland, Umweltbundesamt, Januar 2006

Stahl, Bachmann, Barton, Clark, deFur, Ells, Pittinger, Slimak, Wentsel 2001. Risk Management: Ecological Risk-Based Decision-Making, SETAC Press, 202 pp