



# Global Monitoring Plan data reporting and visualization tool

www.pops-gmp.org

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# The Glpbal Monitoring Plan: data management and visualization tool

# The goal is to

- enhance visibility of the GMP and relevant data,
- facilitate improved interpretation, spatial visualization, and modeling of available monitoring data,
- improve the flow of relevant data to the environmental and health communities,
- enhance our understanding of environmental factors affecting human health and well-being,
- promote a focus on prevention of environmental and human impacts.







# Background - purpose and objective

#### **GMP** for effectiveness evaluation

# **Objective**

Provide a harmonized organizational framework for the collection of comparable monitoring data on the presence of the persistent organic pollutants listed in Annexes A, B and C of the Convention in order to identify time trends in emissions and/or exposure to chemicals in Stockholm Convention as well as to provide information on their regional and global environmental transport.







## GMP - where are we now?

## **GMP** steps/tools

Data generation: monitoring programmes

Data handling: data collection, transfer and storage

Data analysis: analytical tools and reporting



Effectiveness evaluation:

trend assessment - emissions and human exposure evaluation of environmental transport, preparation of regional/global reports for the decision making by Parties at COP7

#### Where are we?

1st report to GMP endorsed at COP4 in 2009 as 5 GMP regional reports in .pdf Effectiveness evaluation - can use the 1st GMP report for baseline, 2nd monitoring report - data collection in 2013/2014, regional reports due in 2015 (COP7)

= need to prepare now for effective linking with previous work and bridging the identified challenges







# GMP - chapter VI - data handling

# What are the building stones?

GMP Guidance - Chapter VI: Data handling

#### Ensure that collected data are

- 1. Relevant and up-to-date
- 2. Have sufficient quality and level of detail
- 3. Consistent and comparable over time
- 4. Transparent and to the extent possible public or unrestricted as much as possible

## 1. Data relevance/scope

- POPs in Annexes A,B,C
  - individual chemicals or cogeners = GMP recommended analytes
- Selected core matrices
  - air, human tissues + new POPs:expanded to water, technical samples







# **Chapter VI**

#### 2. Data structure

Primary GMP data - un-aggregated values; LOD/LOQ use GMP meta-data - complementary information to primary data

- sampling location(s) / site description;
- time of sampling (or the time period represented by the dataset);
- Data on other factors (i.e age/size of animals sampled, volumes of air sampled, information on smoking or dietary habits of the sampled populations, methods employed....);
- Data on parameters to allow conversion between reporting basis (e.g. % lipid and methods used for lipid determination);
- Information on methodologies employed for sampling and analysis,
- Information on QA/QC routines;

Aggregated data: need clear indication of aggregation type: average, geometric mean, median..

+ variability measures (standard deviation, confidence interval etc.)

Uniform methodology for derived parameters - TEQ Application of QA/QC routines







# **Chapter VI**

## 3. Data Quality

Quality check cycle

various levels: at the source, nationally/monitoring programme, regionally same for all regions (not the lowest common denominator)

- use of appropriate methodology and QA/QC routines
- correctness and completeness check regarding the reporting requirements
- data suitability for the purposes of the effectiveness evaluation assessment of data, confidence intervals, supporting information, sampling and analytical methods
- ownership/transparency:
  - Confirmation of correct and accurate transformation/transfer of data
  - Recognize data owners/potential restrictions
  - Require comments from data sources to the output







# Chapter VI - data management

What is the next step for better/most effective use of existing data? How best assist ROGs in their task?

# Chapter VI suggests

Creation of GMP data repository/ regional nodes = 6 geographic regions

- compile and archive national/regional data and outputs
- Facilitate access to data by regional assessment groups
- Ensure transparency, interoperability (forward looking)
- Standardized data exchange and reporting system (comparability, √errors, links)

### Priority tasks/identified challenges

- Based on experience with GMP1 need for standardized data exchange and reporting system what is practical and feasible? How to make it comprehensive AND simple and user friendly?
- Definition of tools for analyses such as time trends + minimum detectable change
- Comprehensive GMP data management e-tool to store, visualize and assess data and assist ROG/GCG most effectively







# First monitoring report

## Chemicals in the Stockholm Convention

### Chemicals identified in GMP reports (isomers, sums, transformation products)

**Aldrin** 

Chlordane

**DDT** 

Dieldrin

Endrin

Heptachlor

Toxaphene

Chlordecone Hexabrombiphenyl

Polychlorinated biphenyls (PCB)

dibenzofurans (PCDDs/PCDFs)

Tetrabromodiphenyl ether (tetraBDE) Pentabromodiphenyl ether (pentaBDE)

and perfluoroctane sulfonyl fluoride

Heptabfomodiphenyl ether (hepta BDE) Hexabromdiphenyl ether (hexa BDE)

Perfluorooctane sulfonic acid (PFOA), its salts

Alpha hexachlorocyclohexane (a-HCH)\* Beta hexachlorocyclohexane (b-HCH)\*

Pentachlorbenzene (PeCB)

Polychlorinated dibenzo-p-dioxins and

**Aldrin** 

Cis- (alpha-) chlordane, Trans- (gamma-) chlordane, Oxychlordane, Cis- and Trans-nonachlor, Chlordane (as a group), Chlordane + Nonachlor, Chlordane + trans-nonachlor

DDT, DDE, DDD, o,p'-DDT, o,p'-DDE, o,p'-DDD, p,p'-DDT, p,p'-DDE, p,p'-DDD,

Sum of DDTs, Sum p,p-DDX (p,p-isomers together), Sums of various number of DDT isomers, DDT + p,p-DDE

**Dieldrin, Endrin** (also **ketone** and Endrin group = sum of)

Heptachlor, Heptachlorepoxide, Heptachlorepoxide cis-, Heptachlorepoxide trans-, Heptachlor group (sum of)

**Hexachlorbenzene (HCB)** 

Hexachlorbenzene (HCB)

Mirex Mirex

> 36 various PCBs: indicator PCBs (PCB 28, PCB 52, PCB 101, PCB 118, PCB 138, PCB 153, PCB 180) and 29 others. Various PCBs WHO-TEQ and PCBs I-TEQ, Sum of 3, 10, 14, 21, 25, 48 PCBs, dI-PCBs (also with WHO-TEQ), Mono-ortho and Non-ortho PCBs in TEQs

> PCDDs and PCDFs (15 congeners) in sum or separately - PCDDs, PCDFs, PCDDs/Fs - often as various TEQs: WHO-TEQ, Nordic-TEQ and I-TEQ, OCDD, OCDF

Parlar (Toxaphene) 26 isomers, Parlar (Toxaphene) 50, Parlar (Toxaphene) 62, Toxaphene (as a group), Parlar 40, Parlar 41. Parlar 44

Pentachlorbenzene (PeCB)

Reported only as a sum of 3 PBDEs

a-HCH, b-HCH, g-HCH, d-HCH, HCHs (sum of)

Endosulfan

Lindane (g-HCH)\*

Endosulfan I, Endosulfan II, Endosulfan SO4, Endosulfans (sum of)

+ other chemicals not listed in the Stockholm Convention Dacthal, Isodine, Methoxychlor, trifluralin and delta-HCH





## Parameters in GMR1

Together, all regional GMP reports contain 171 variables (including concentration data on congeners, isomers, transformation products, various summations and toxic equivalents – TEQs). Analysing the primary pool of reported parameters, 58 of them were related to 12 original Stockholm Convention POPs) as specified in the chapter 2 of the *Guidance on the GMP for Persistent Organic Pollutants, 2007* (compounds highly recommended for monitoring and evaluation).

Additional 7 of all reported variables (alpha-HCH, beta-HCH, gamma-HCH, PeCB, endosulfan I, endosulfan II and endosulfan SO<sub>4</sub>) were not obligatory at the time of the first GMP report, but they are very important for the future as they are related to the additional 10 compounds that were listed in the Stockholm Convention in 2009 and in 2011, respectively.

Their specification can be found in the revised *Guidance on the GMP for Persistent Organic Pollutants*, 2009. Draft revised guidance on the global monitoring plan for persistent organic pollutants.







### Parameters in GMR1

The third group (84 variables) comprises parameters related to the Stockholm Convention POPs, but not specified in the Guidelines. Among them, there are various PCB congeners (not recommended in the Guidance) and often not very well defined summations of the individual compounds (56 together) as well as variety of 28 toxic equivalents (TEQs for dioxins, furans and PCBs based on various TEFs, often not cited correctly).

In the fourth group there are 22 chemicals or parameters with no relation to the Stockholm Convention (mostly PAHs (16 PAHs + sum of PAHs). We believe that only the first two groups of parameters should be reported in the next GMP data collection campaigns as parameters from the third group can be calculated in the (global) database and parameters currently reported under the fourth group have no relation to the effectiveness evaluation of the Stockholm Convention.







## Identified data challenges

- heterogenous nomenclature (i.e. trans-chlordane vs. gammachlordane)
- unclear reporting or missing units
- parameters without specification which isomers or degradation products summed up ("chlordanes", "PCBs", "DDTs", "heptachlor)
- Difference in labeling of PCBs summations
- Limit of Quantification (LOQ) not reported
- Ambient air data reported as spatially aggregated values, often without specific information on number and type of sampling site
- Missing or wrong specification of the TEF values used for calculation of the TEQ (year).
- Various WHO-TEQ values used (WHO-TEQ 1995, 1997, 1998, 2001, 2005)

## Consequence

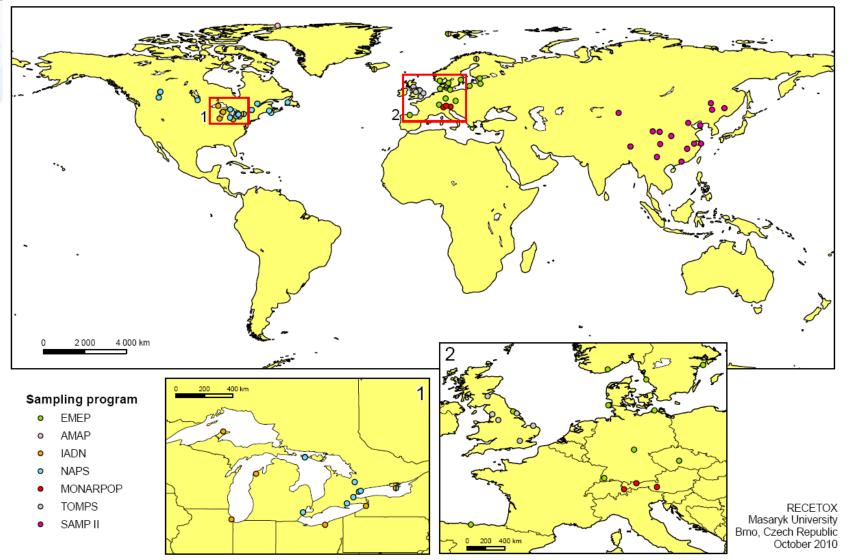
- reliability of collected GMP1 data significantly influenced
- narrowed data pool for future comparisons BUT could be improved if we can find the way how to amend/complement problematic records in GMP1 and effectively link them to GMP2







# Long-term active air sampling networks

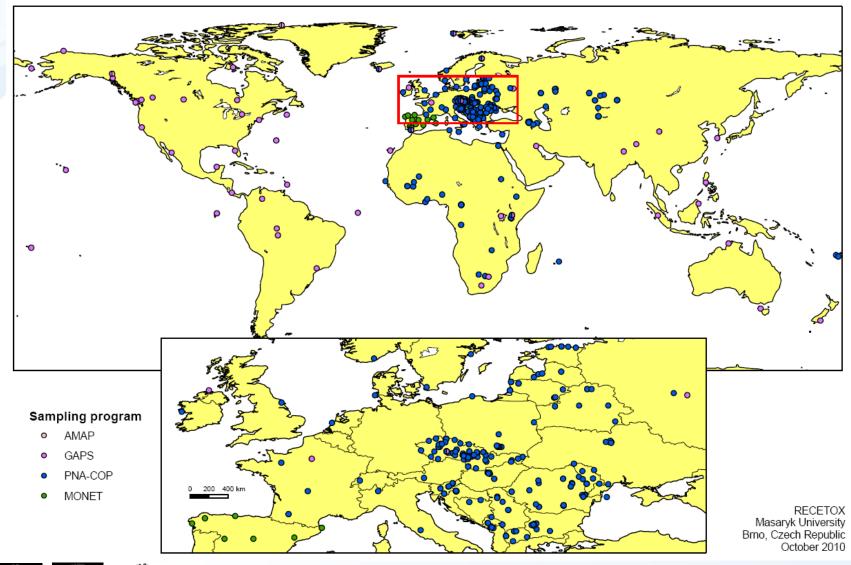








# Passive air sampling networks









Click on this logo to

return to this page again



WORLD MAP -MONITORING OVERVIEW SAMPLING FREQUENCY -PARAMETERS SAMPLING FREQUENCY -YEARS

REPORTED VALUES

Main menu

#### WORLD MAP – MONITORING OVERVIEW

An interactive world map displaying monitoring activities within all worlds' countries in individual years.

» Open the visualisation

#### SAMPLING FREQUENCY – PARAMETERS

The chart shows sampling frequency of individual compounds in countries.

» Open the visualisation

Description and links to all available reports

#### SAMPLING FREQUENCY – YEARS

The chart shows sampling frequency of individual compounds on 1-year divided timeline.

» Open the visualisation

# REPORTED VALUES

This chart is designed for direct presentation of reported values and concentrations.

» Open the visualisation









#### **GMP Reports On-line Data Visualzation**

Back to home page



WORLD MAP -MONITORING OVERVIEW

SAMPLING FREQUENCY -**PARAMETERS** 

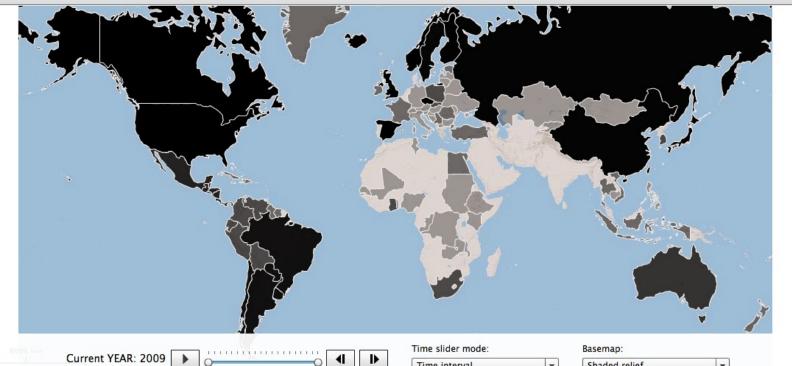
SAMPLING FREQUENCY -YEARS

REPORTED VALUES



#### WORLD MAP - MONITORING OVERVIEW

Matrix: Air





Time interval

Shaded relief







### **SAMPLING FREQUENCY – YEARS**



Matrix:	UN region:	Parameter:	Year (1960 - 2010):	
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USA - Barrow				
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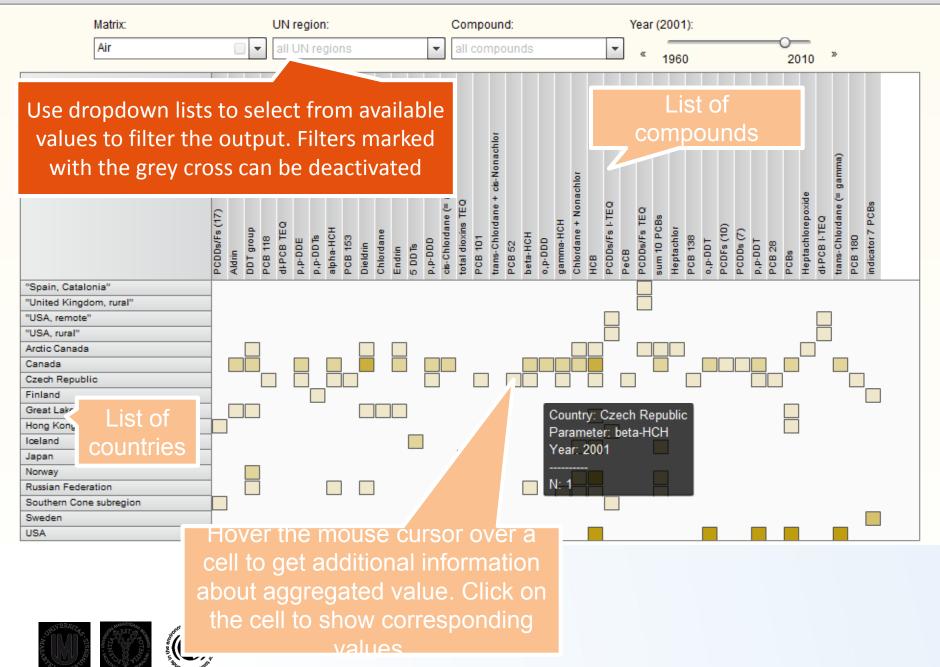






#### SAMPLING FREQUENCY – PARAMETERS





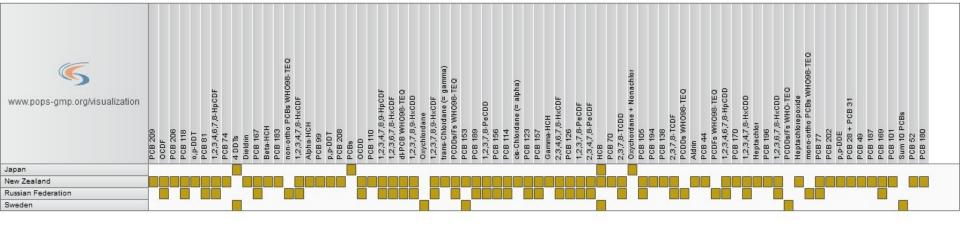
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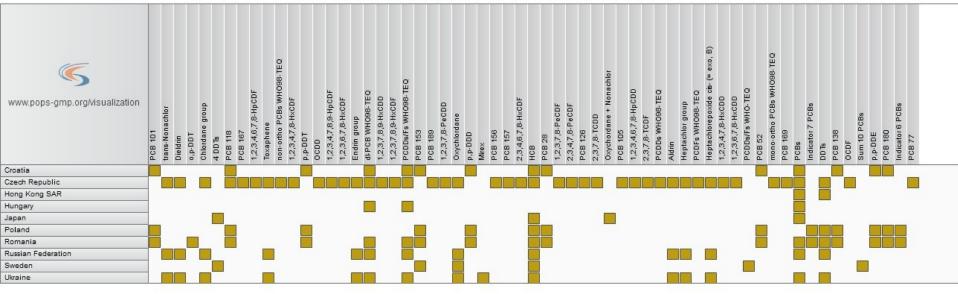








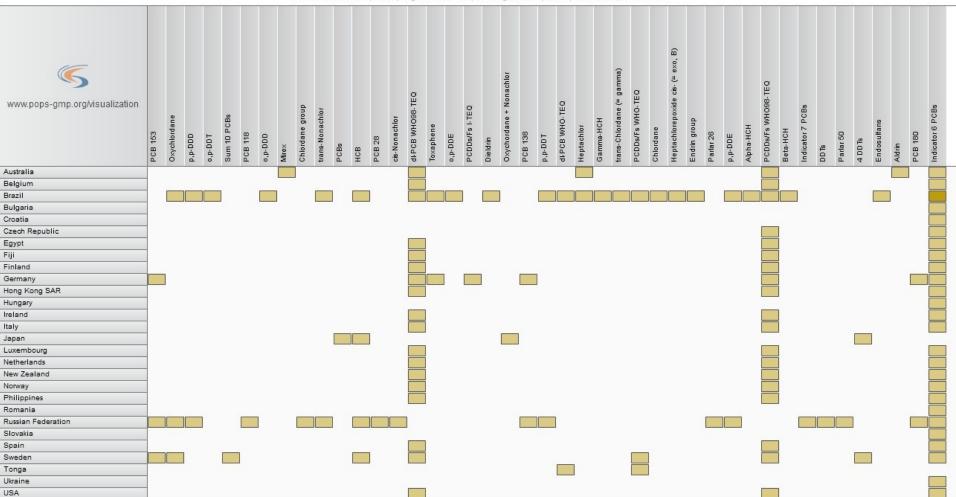








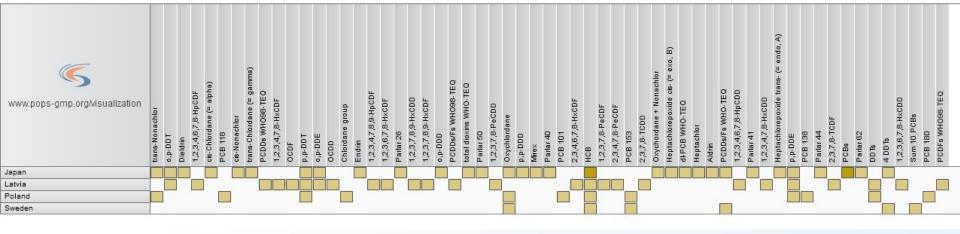








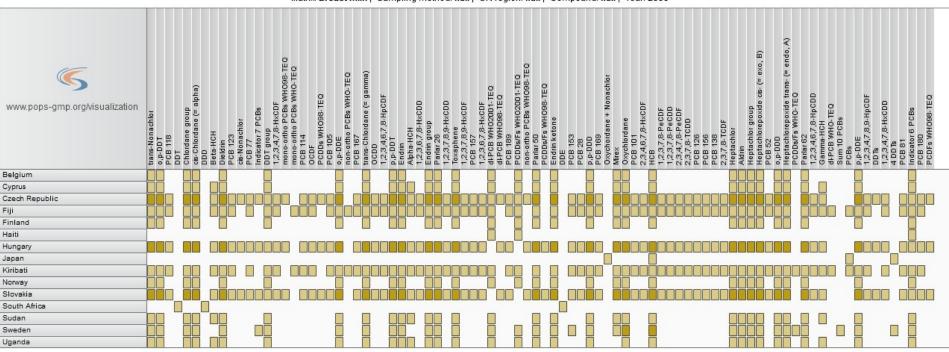


















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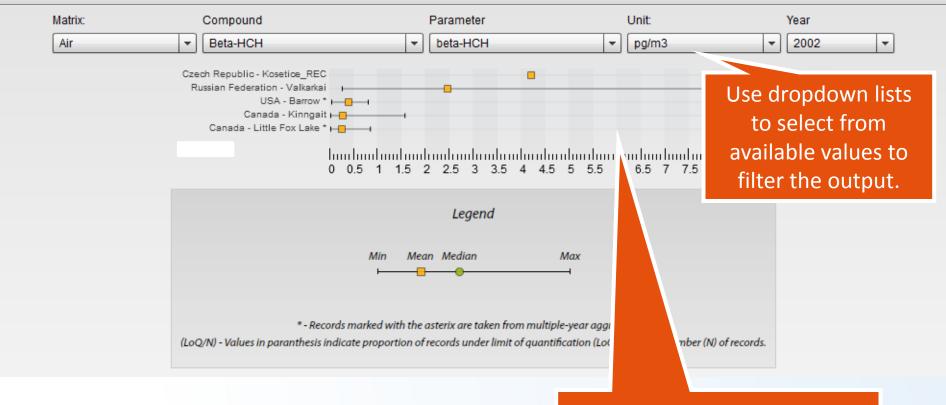






#### REPORTED VALUES





Reported values are shown as a box-and-whisker plot. A plot consist of minimum, maximum, mean and median.

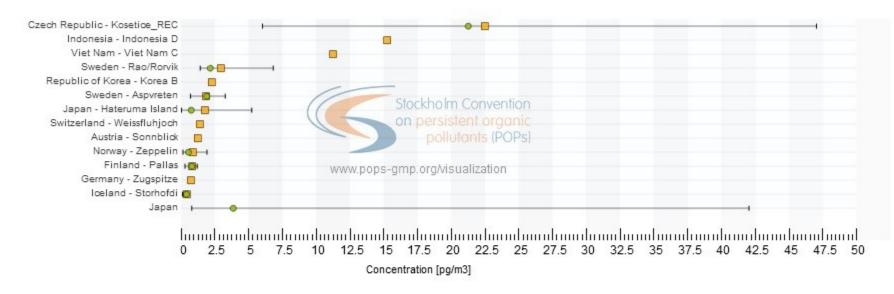
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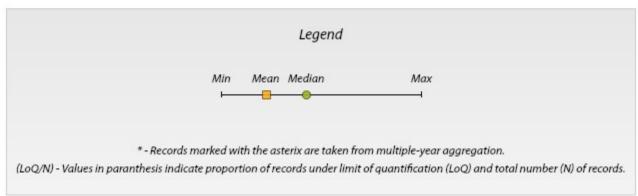






Matrix: Air | Sampling method: active | Compound: Alpha-HCH | Parameter: Alpha-HCH | Unit: pg/m3 | Year: 2005



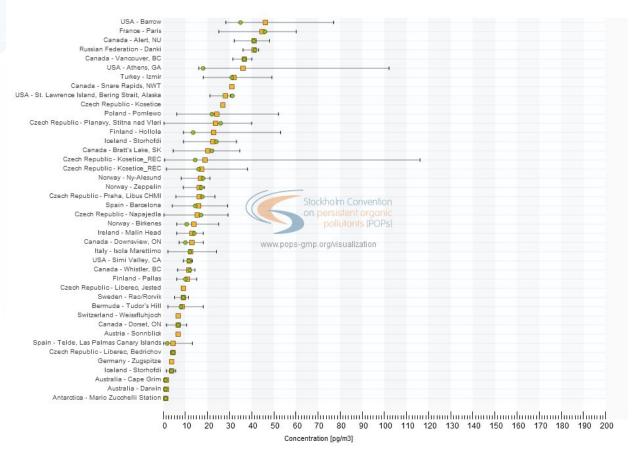


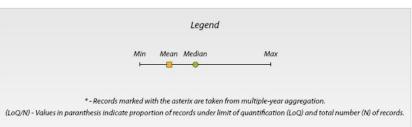






Matrix: Air | Sampling method: active & passive | Compound: Alpha-HCH | Parameter: Alpha-HCH | Unit: pg/m3 | Year: 2005



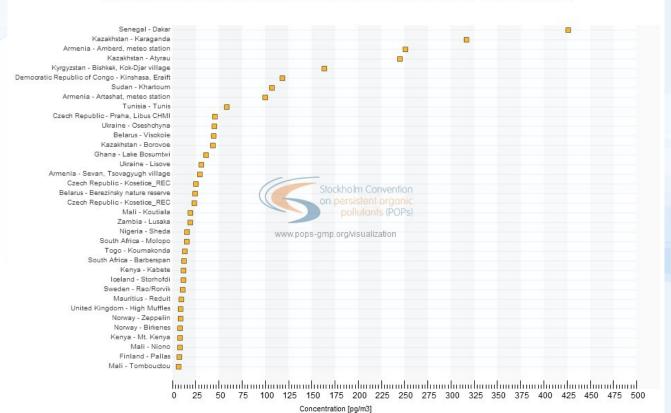








Matrix: Air | Sampling method: active & passive | Compound: PCB | Parameter: Indicator 6 PCBs | Unit: pg/m3 | Year: 2008



Legend

Min Mean Median Max

\*-Records marked with the asterix are taken from multiple-year aggregation.

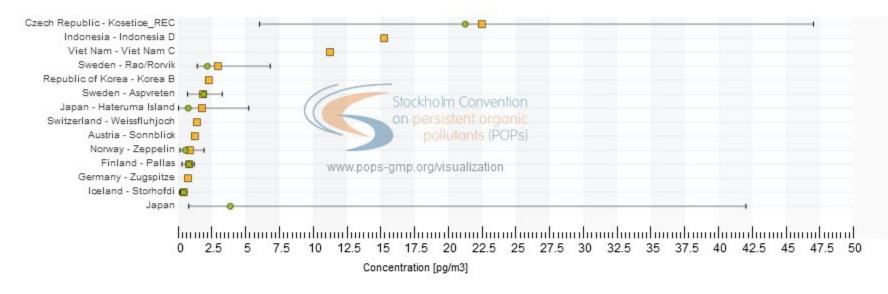
(LoQ/N) - Values in paranthesis indicate proportion of records under limit of quantification (LoQ) and total number (N) of records.

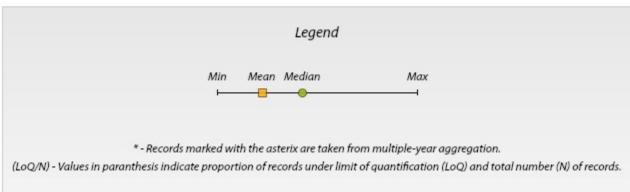






Matrix: Air | Sampling method: active | Compound: DDT | Parameter: p,p-DDE | Unit: pg/m3 | Year: 2005



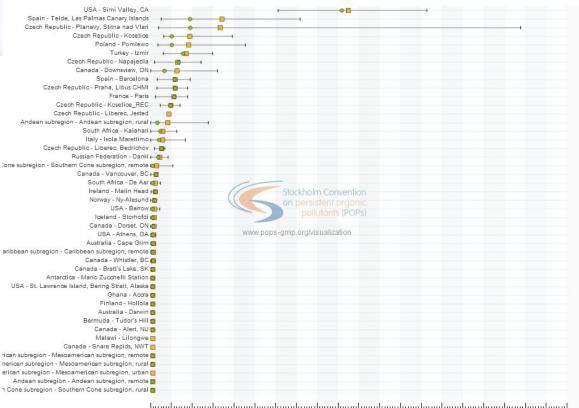




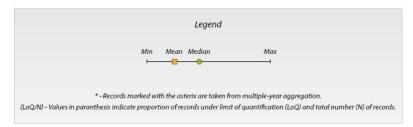




Matrix: Air | Sampling method: passive | Compound: DDT | Parameter: p,p-DDE | Unit: pg/m3 | Year: 2005



| 10 | 35 | 70 | 105 | 140 | 175 | 210 | 245 | 280 | 315 | 350 | 385 | 420 | 455 | 490 | 525 | 560 | 595 | 630 | 665 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 700 | 7

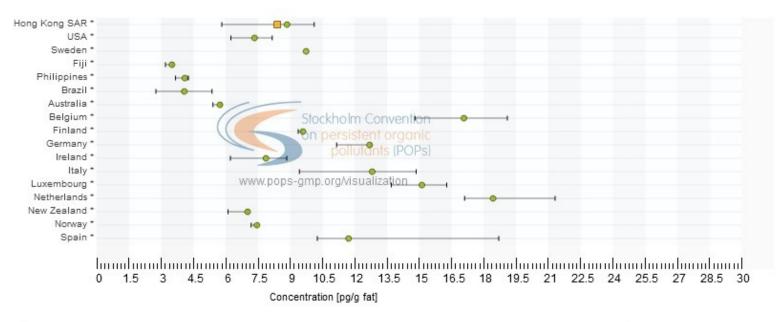


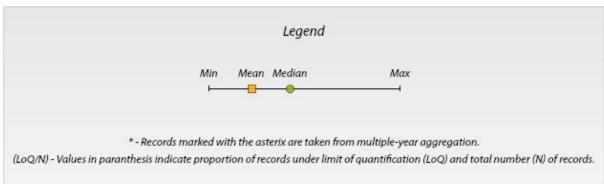






Matrix: Breast milk | Sampling method: null | Compound: PCDF | Parameter: PCDDs/Fs WHO98-TEQ | Unit: pg/g fat | Year: 2002



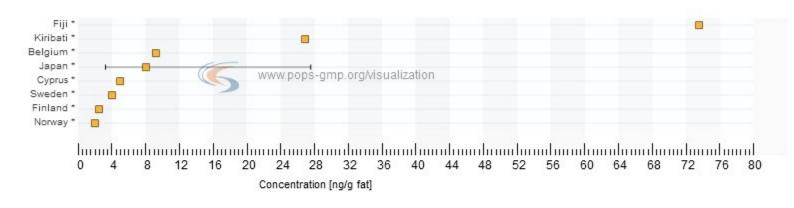


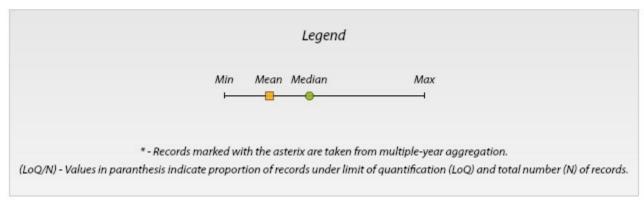






Matrix: Breast milk | Sampling method: null | Compound: DDT | Parameter: p,p-DDT | Unit: ng/g fat | Year: 2005

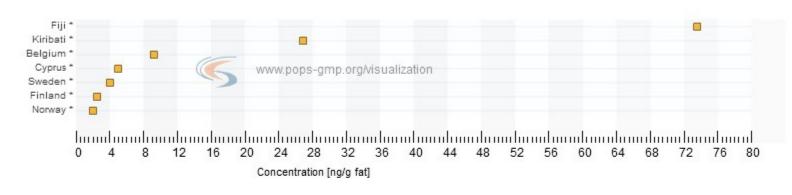


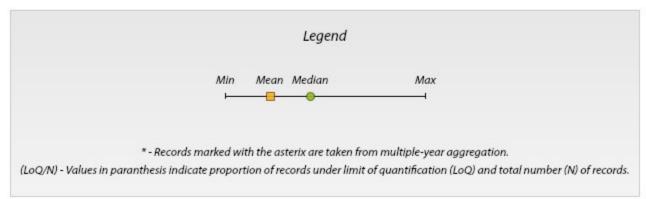








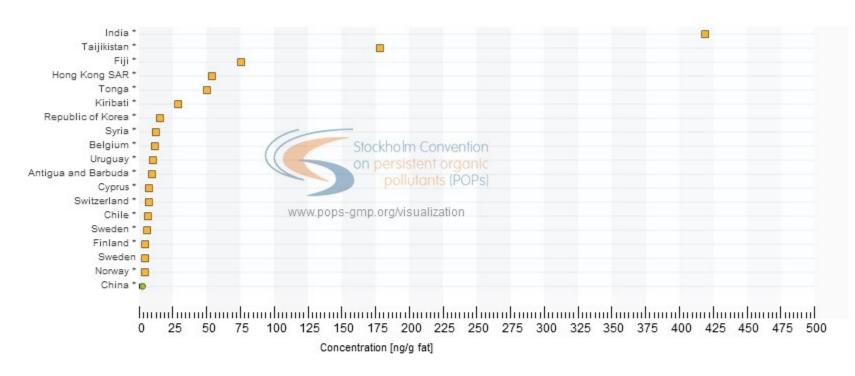


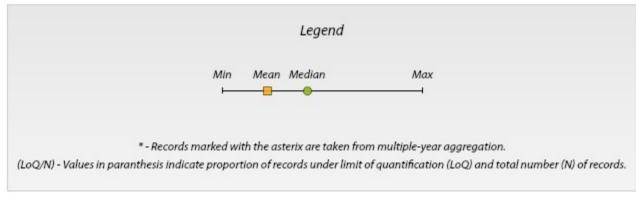


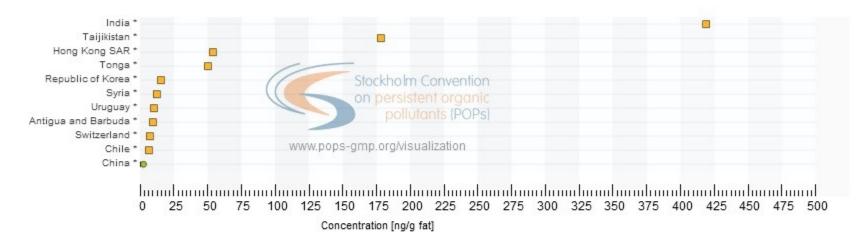


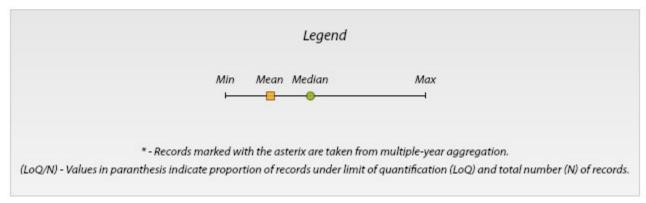








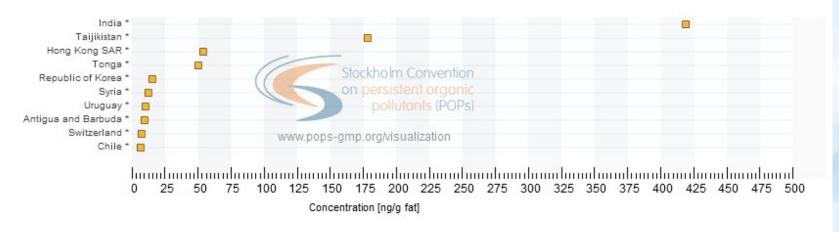


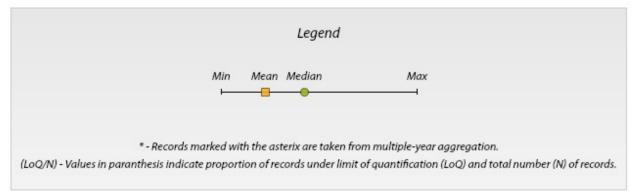










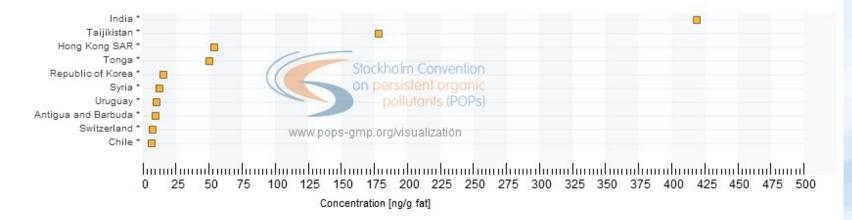








Matrix: Breast milk | Sampling method: null | Compound: DDT | Parameter: p,p-DDT | Unit: ng/g fat | Year: 2010





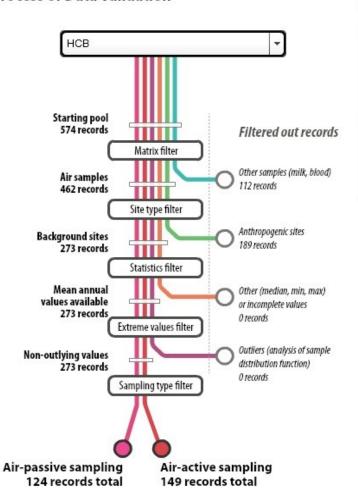




### GMP Reports On-line Data Visualization REGIONAL BACKGROUNDS - data validation

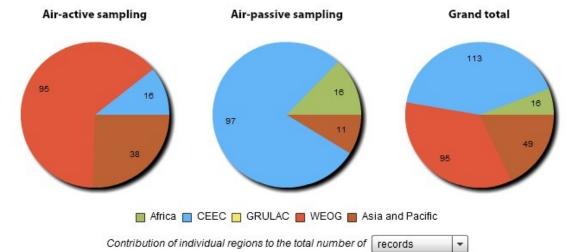
Parameter: HCB |

#### **Process of Data Validation**



#### **Results of Data Validation**

Sampling method	Africa	CEEC	GRULAC	WEOG	Asia and P	TOTAL
HCB Air-passive sampling	12 countries 16 sites 16 records 2008 - 2008	19 countries 66 sites 97 records 2003 - 2008	0 countries 0 sites 0 records	0 countries 0 sites 0 records	3 countries 8 sites 11 records 2006 - 2008	34 countries 90 sites 124 records 2003 - 2008
HCB Air-active sampling	0 countries 0 sites 0 records	2 countries 3 sites 16 records 1998 - 2008	0 countries 0 sites 0 records	8 countries 19 sites 95 records 1998 - 2008	9 countries 32 sites 38 records 2004 - 2007	19 countries 54 sites 149 records 1998 - 2008

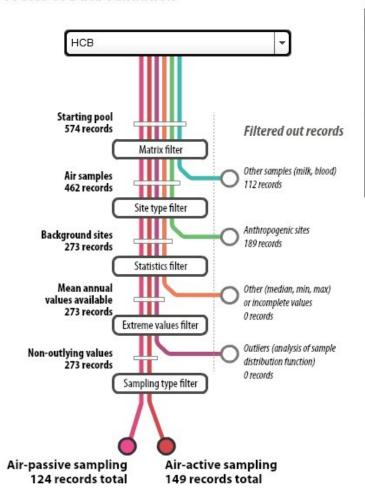




### GMP Reports On-line Data Visualization REGIONAL BACKGROUNDS - data validation

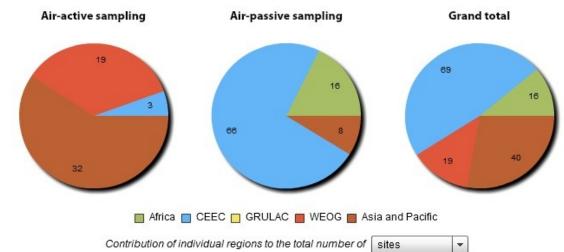
Parameter: HCB |

#### **Process of Data Validation**



#### **Results of Data Validation**

Sampling method	Africa	CEEC	GRULAC	WEOG	Asia and P	TOTAL
HCB Air-passive sampling	12 countries 16 sites 16 records 2008 - 2008	19 countries 66 sites 97 records 2003 - 2008	0 countries 0 sites 0 records	0 countries 0 sites 0 records	<ul><li>3 countries</li><li>8 sites</li><li>11 records</li><li>2006 - 2008</li></ul>	34 countries 90 sites 124 records 2003 - 2008
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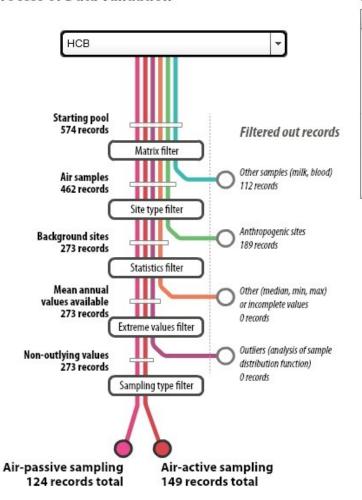




### GMP Reports On-line Data Visualization REGIONAL BACKGROUNDS - data validation

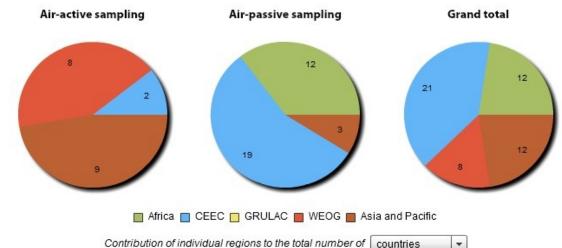
Parameter: HCB |

#### **Process of Data Validation**



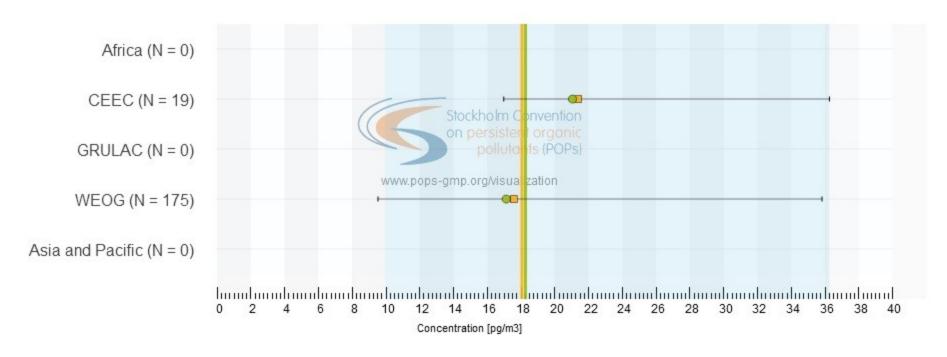
#### **Results of Data Validation**

Sampling method	Africa	CEEC	GRULAC	WEOG	Asia and P	TOTAL
HCB Air-passive sampling	12 countries 16 sites 16 records 2008 - 2008	19 countries 66 sites 97 records 2003 - 2008	0 countries 0 sites 0 records	0 countries 0 sites 0 records	<ul><li>3 countries</li><li>8 sites</li><li>11 records</li><li>2006 - 2008</li></ul>	34 countries 90 sites 124 records 2003 - 2008
HCB Air-active sampling	0 countries 0 sites 0 records	2 countries 3 sites 16 records 1998 - 2008	0 countries 0 sites 0 records	8 countries 19 sites 95 records 1998 - 2008	9 countries 32 sites 38 records 2004 - 2007	19 countries 54 sites 149 records 1998 - 2008





Parameter: Alpha-HCH | Sampling method: active

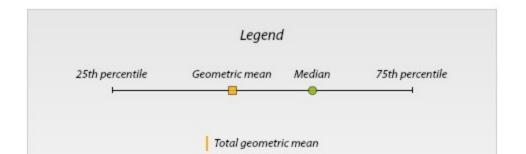


Total median: 18.1 pg/m3

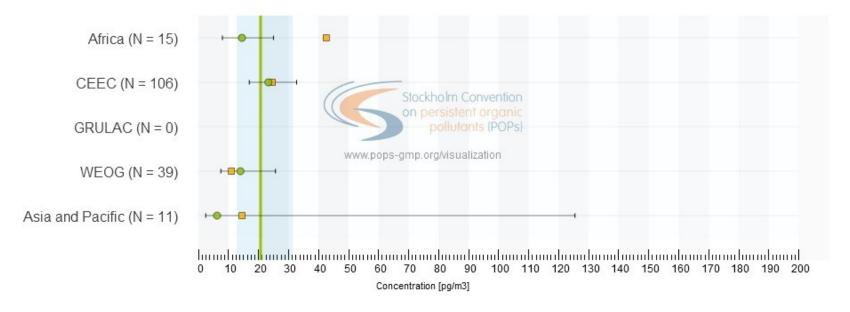
Total geometric mean: 17.897 pg/m3

Total 25th percentile: 9.862 pg/m3

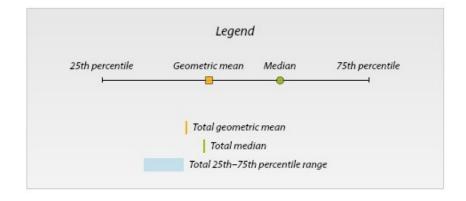
Total 75th percentile: 35.853 pg/m3



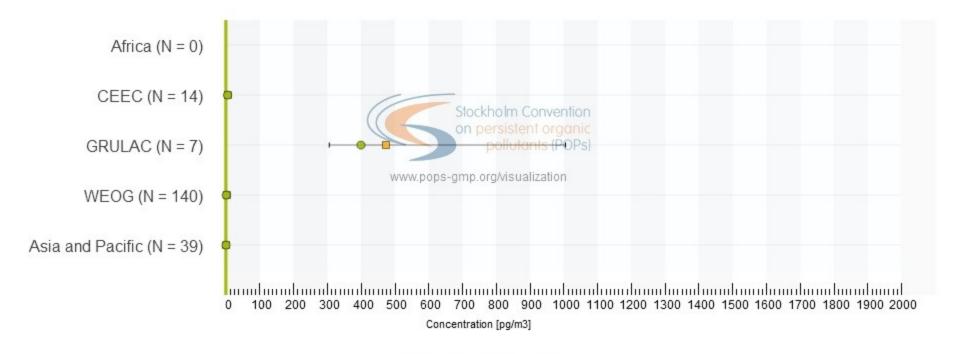
Parameter: Alpha-HCH | Sampling method: passive



Total median: 20.854 pg/m3 Total geometric mean: 20.899 pg/m3 Total 25th percentile: 12.786 pg/m3 Total 75th percentile: 31.495 pg/m3



Parameter: p,p-DDT | Sampling method: active

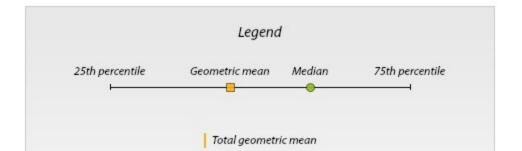


Total median: 0.796 pg/m3

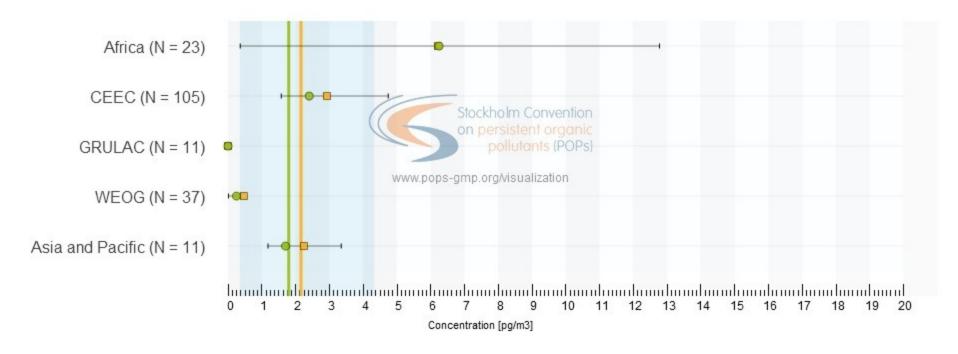
Total geometric mean: 1.76 pg/m3

Total 25th percentile: 0.25 pg/m3

Total 75th percentile: 2.596 pg/m3



Parameter: p,p-DDT | Sampling method: passive

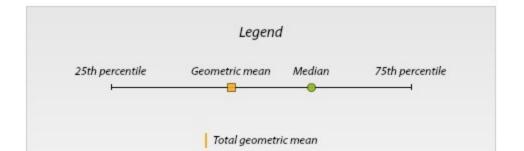


Total median: 1.813 pg/m3

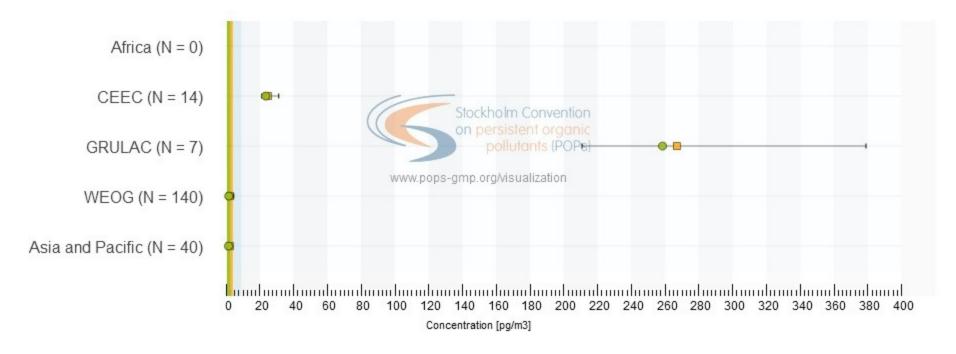
Total geometric mean: 2.173 pg/m3

Total 25th percentile: 0.35 pg/m3

Total 75th percentile: 4.33 pg/m3



Parameter: p,p-DDE | Sampling method: active

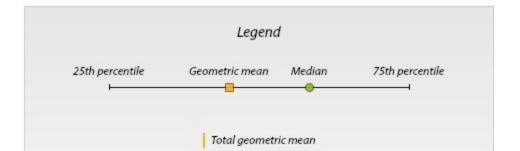


Total median: 1.823 pg/m3

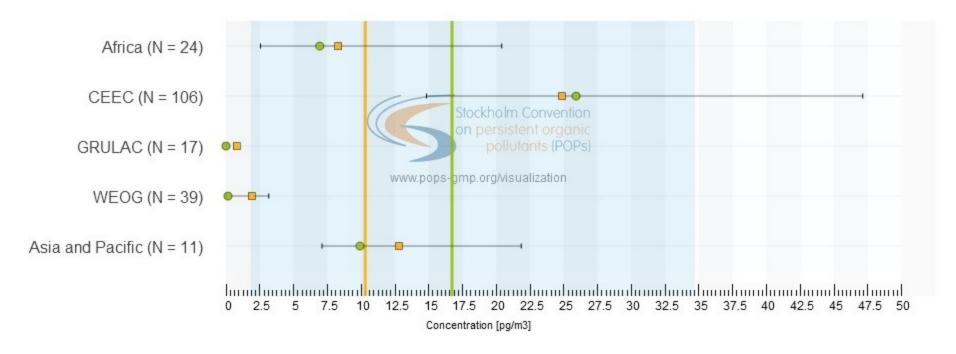
Total geometric mean: 3.477 pg/m3

Total 25th percentile: 0.74 pg/m3

Total 75th percentile: 8.99 pg/m3



Parameter: p,p-DDE | Sampling method: passive

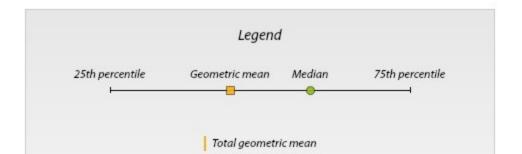


Total median: 16.763 pg/m3

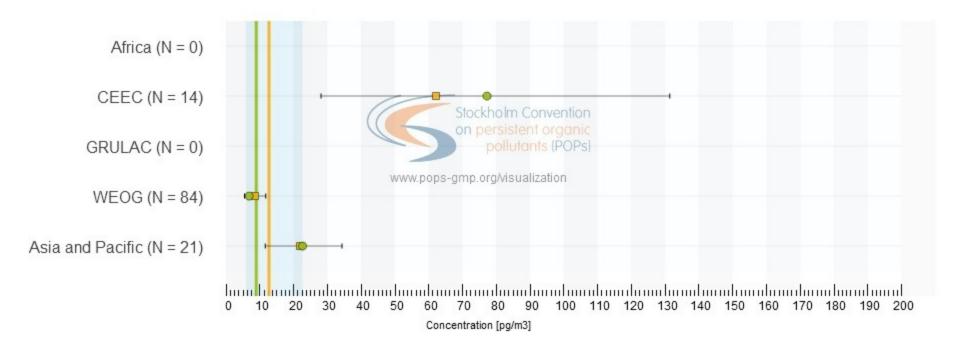
Total geometric mean: 10.337 pg/m3

Total 25th percentile: 1.84 pg/m3

Total 75th percentile: 34.7 pg/m3



Parameter: Indicator 6 PCBs | Sampling method: active

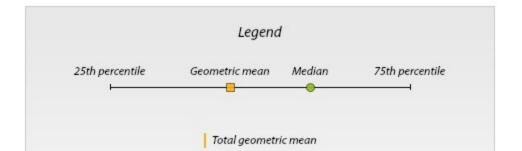


Total median: 9.0746 pg/m3

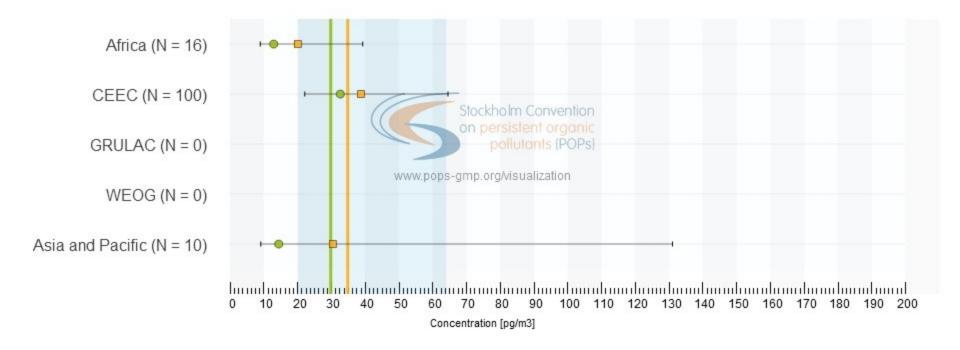
Total geometric mean: 12.896 pg/m3

Total 25th percentile: 5.915 pg/m3

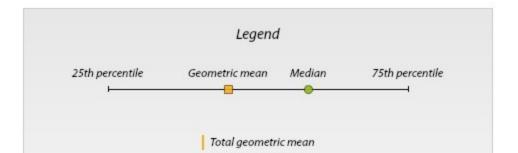
Total 75th percentile: 22.664 pg/m3



Parameter: Indicator 6 PCBs | Sampling method: passive

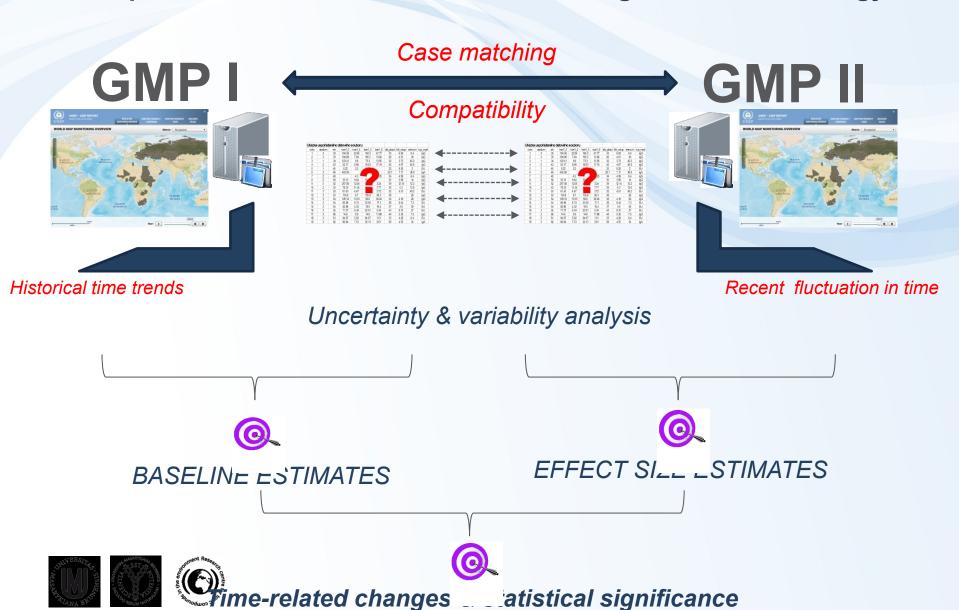


Total median: 29.975 pg/m3 Total geometric mean: 34.991 pg/m3 Total 25th percentile: 20.065 pg/m3 Total 75th percentile: 64.0575 pg/m3



# Data analysis - methodological proposal

3. Steps for standardization of the monitoring data - methodology







Retrospective corrections

ı.

NOMENCLATURE

П.

**CLASSIFIERS** 

ш.

**INFORMATION SOURCE IDENTIFIERS** 

IV.

**OBLIGATORY DESCRIPTORS** 





Real time validatoin



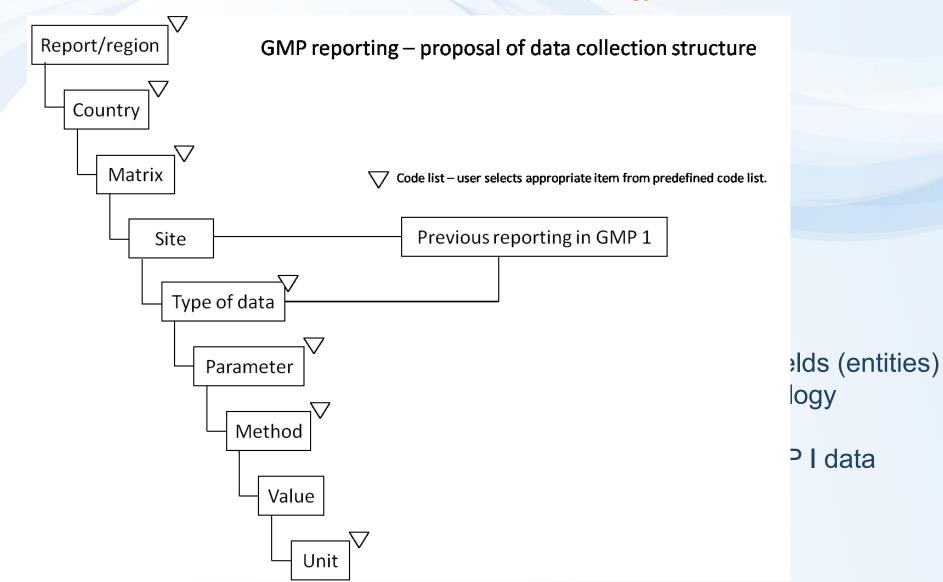




# Proposal for e-data capture system

#### 4. Proposal for electronic format template for the next GMP

Hierarchical structure of data entities and umbrella of ontology



#### Proposal for electronic reporting format - overview

- Ensures that GMP 2 could be effectively linked to GMP1 reports
- Data are relevant (and up-to-date) SC chemicals only
- Data have sufficient quality and level of detail analytical tools
- Consistent and comparable over time hierarchical structure and optimized ontology
- Access is transparent via web interface,
- data restriction possible to set up depending on the user/provider/data owner
- Safety and security of data international standards and settings in place
- Simple and feasible step-by-step approach to GMP data storage and handling, including data search and data analysis tools - now available for trials/testing













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# Thank you for your kind attention





