

Hoe bang moeten we zijn voor bestrijdingsmiddelen?

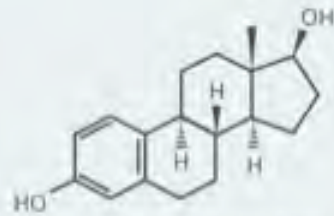
Martin van den Berg

Institute for Risk Assessment Sciences (IRAS)

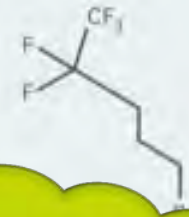
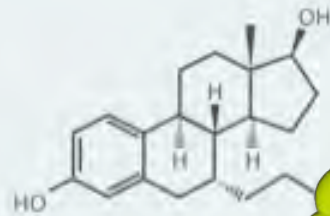
Universiteit Utrecht

m.vandenberg@uu.nl



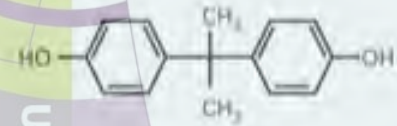


E₂ (17β-estradiol)

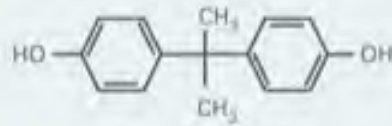


De spelregels van de toxicologie

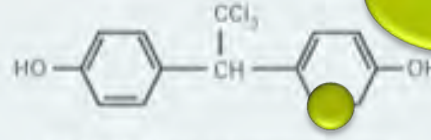
Group 1



BPA (bisphenol A)



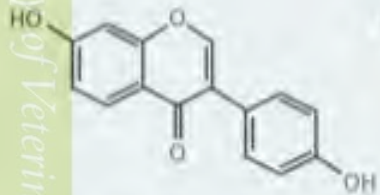
BPAF (bisphenol AF)



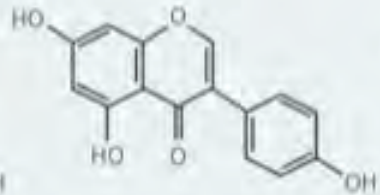
HPTE [2,2-bis(p-hydroxyphenyl)-chloroethane]

4n-NP (4-n-nonylphenol)

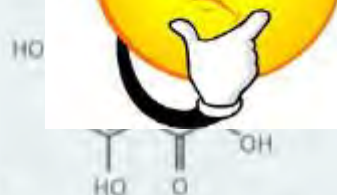
Group 2



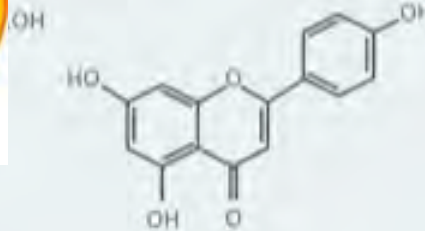
Dai (daidzein)



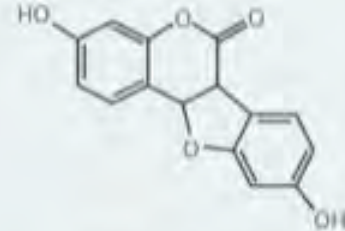
Gen (genistein)



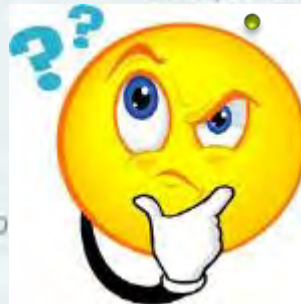
Kaem (kaempferol)



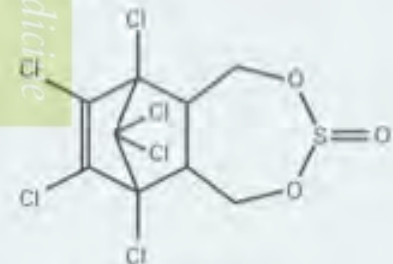
Api (apigenin)



Coum (coumestrol)



Group 3



Endo (endosulfan)



Kep (kepone)



1-BP (1-bromopropane)

Hazard vs. Risk

Universiteit Utrecht

- **Hazard**

(potentieel gevaarlijk voor gezondheid):

= intrinsieke eigenschap van een stof

- **Risk (risico):**

= waarschijnlijkheid dat die gevaarlijke eigenschap zich manifesteert bij een bepaalde blootstelling

Group 1

UU

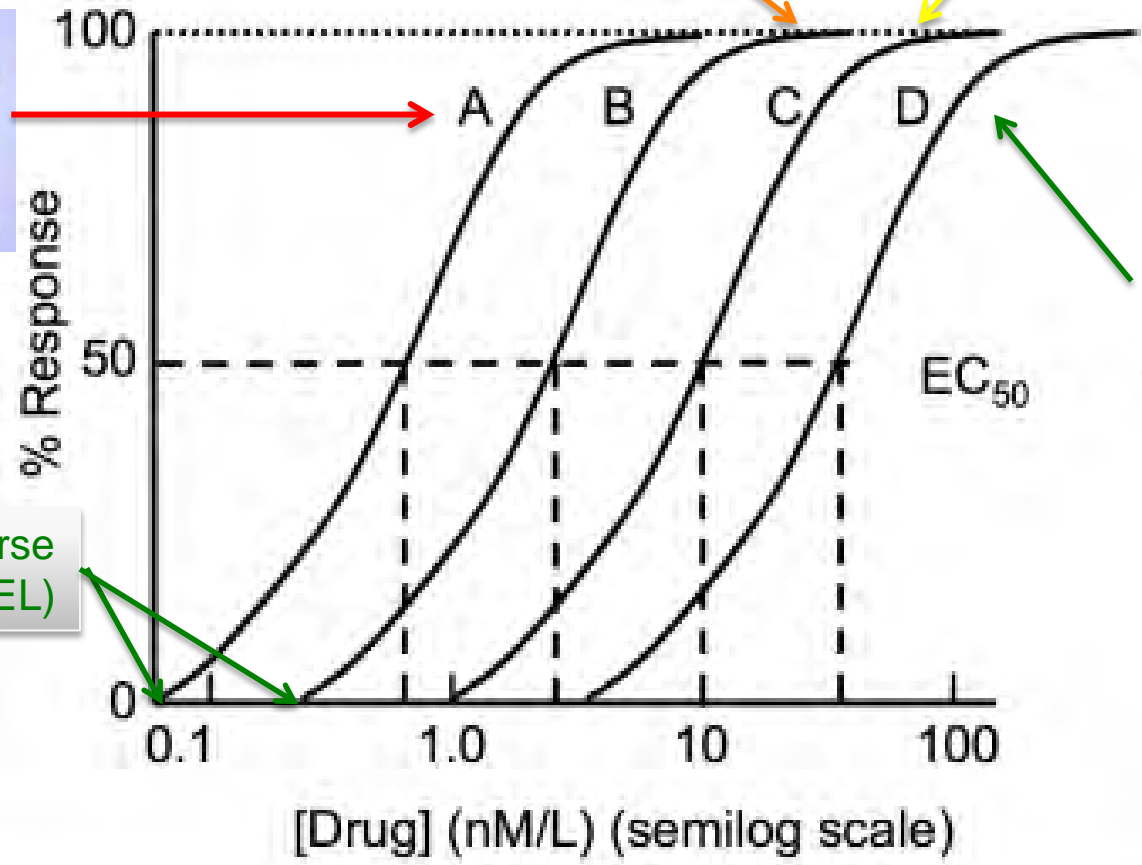
Faculty of Veterinary Medicine





Risico schatting in de toxicologie voor de mens

1. Identificatie gevaarlijke eigenschap (hazard)
 2. Karakterisering eigenschap (dosis-effect)
 3. Vaststellen van de blootstelling
 4. Vaststellen van het risico
-
5. Maatregelen om risico te beperken
 6. Risico communiceren



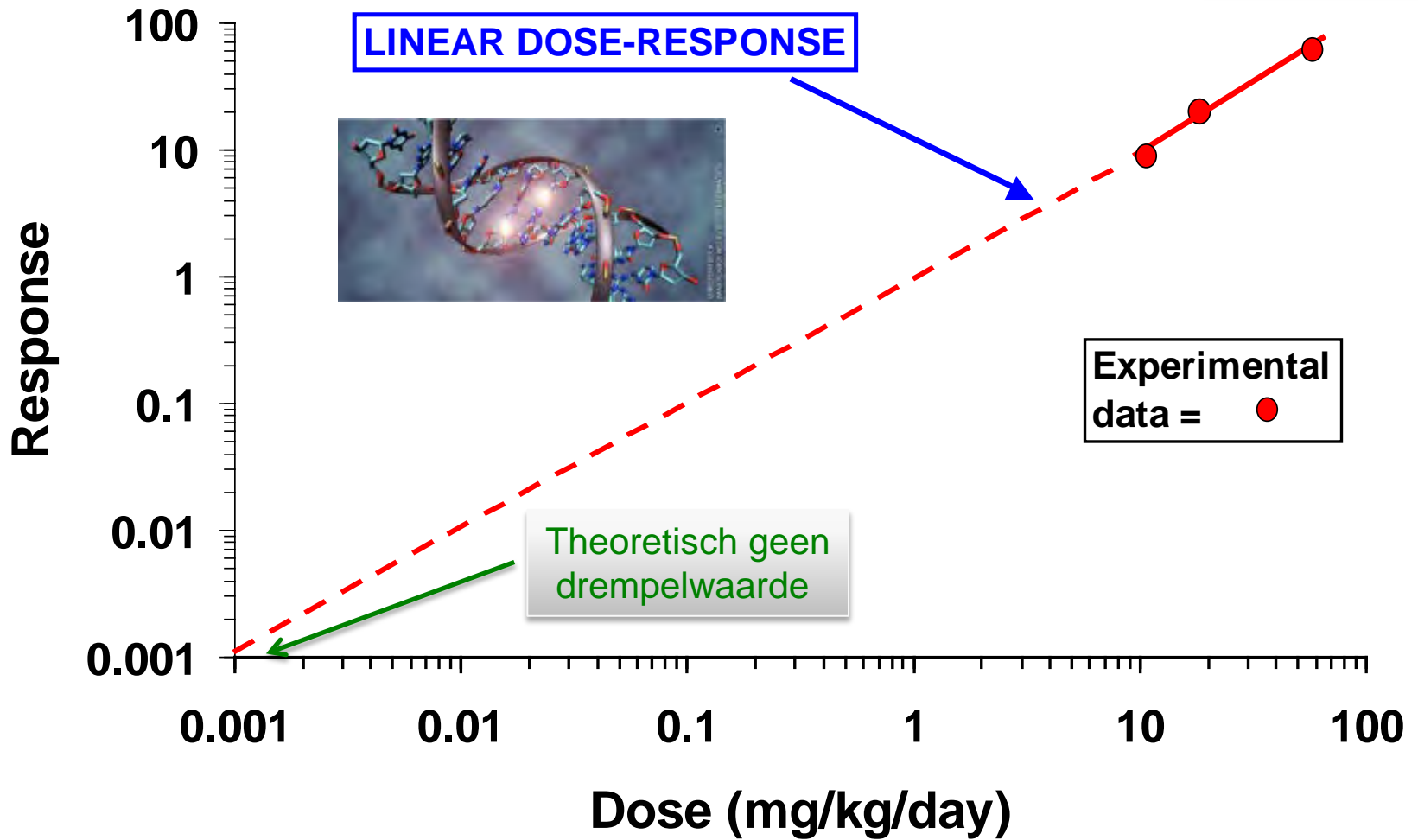
No Observed Adverse Effect Level (NOAEL)

Safety Factor Default 100x

Gevoeligheid voor een stof hangt af van levensfase

Hypothetical dose-response

Bijv. tumoren bij de rat



Acceptabel: 1: 100.00 tot 1.000.000 mensen ontwikkelt een tumor



“The problem with toxicology is not the practicing toxicologists, but chemists who can detect, precisely, toxicologically insignificant amounts of chemicals”

Rene Truhaut, University of Paris (1909-1994)

Detectie grens tegenwoordig: *femtogram* = 10^{-15} gram



News

Carcinogenicity of tetrachlorvinphos, parathion, malathion, diazinon, and glyphosate



In March, 2015, 17 experts from 11 countries met at the International Agency for Research on Cancer (IARC) in Lyon, France to assess the carcinogenicity of the organophosphate pesticides tetrachlorvinphos, parathion, malathion, diazinon, and glyphosate (table). These assessments will be published as volume 112 of the IARC Monographs.¹

The insecticides tetrachlorvinphos and parathion were classified as "possibly carcinogenic to humans" (Group 2B). The evidence from human studies was scarce and considered inadequate. Tetrachlorvinphos induced hepatocellular tumours (benign or malignant) in mice, renal tubule tumours (benign or malignant) in male mice,² and spleen haemangioma in male rats. Tetrachlorvinphos is a reactive oxon with affinity for esterases. In experimental animals, tetrachlorvinphos is systematically eliminated in urine. Although bacterial mutagenesis tests were negative, tetrachlorvinphos induced genotoxicity in some assays (chromosomal damage in rats and in vitro) and increased cell proliferation (hyperplasia in rodents). Tetrachlorvinphos is banned in the European Union. In the USA, it continues to be used on animals, including in pet flea collars.

For parathion, associations with cancers in several tissues were observed in occupational studies, but the evidence in humans remains sparse. In mice, parathion increased bronchioloalveolar adenoma and/or carcinoma in males and lymphoma in females. In rats, parathion induced adrenal cortical adenoma or carcinoma (combined),³ malignant pancreatic tumours, and thyroid follicular cell adenoma in males (after subcutaneous injection) and mammary gland adenocarcinoma (after subcutaneous injection in females).⁴ Parathion is rapidly absorbed and distributed. Parathion metabolism

to the bioactive metabolite, paraoxon, is similar across species. Although bacterial mutagenesis tests were negative, parathion induced DNA and chromosomal damage in human cells in vitro. Parathion markedly increased rat mammary gland terminal end bud density.⁴ Parathion use has been severely restricted since the 1980s.

The insecticides malathion and diazinon were classified as "probably carcinogenic to humans" (Group 2A). Malathion is used in agriculture, public health, and residential insect control. It continues to be produced in substantial volumes throughout the world. There is limited evidence in humans for the carcinogenicity of malathion. Case-exposures reported positive associations with non-Hodgkin lymphoma in the USA,⁵ Canada,⁶ and Sweden,⁷ although no increased risk of non-Hodgkin lymphoma was observed in the large Agricultural Health Study cohort (AHS). Occupational use was associated with an increased risk of prostate cancer in a Canadian case-control study⁸ and in the AHS, which reported a significant trend for

aggressive cancers after adjustment for other pesticides.⁹ In mice, malathion increased hepatocellular adenoma or carcinoma (combined).¹⁰ In rats, it increased thyroid carcinoma in males, hepatocellular adenoma or carcinoma (combined) in females, and mammary gland adenocarcinoma after subcutaneous injection in females.⁴ Malathion is rapidly absorbed and distributed. Metabolism to the bioactive metabolite, malaoxon, is similar across species. Malaoxon strongly inhibits esterases; atropine reduced carcinogenesis-related effects in one study.⁴ Malathion induced DNA and chromosomal damage in humans, corroborated by studies in animals in vitro. Bacterial mutagenesis tests supported disruption of hormone pathways. Hormonal effects probably mediate rodent thyroid and mammary gland proliferation.

Diazinon has been applied in agriculture and for control of home and garden insects. There was limited evidence for diazinon carcinogenicity in humans. Positive associations for non-Hodgkin lymphoma, with

Lennox Oncol 2015
Published Online
March 20, 2015
http://dx.doi.org/10.1016/j.ijom.2015.03.016
51470-2045(15)70134-8
For more on the IARC Monographs see <http://www.monographs.iarc.fr>
Upcoming meetings
June 2-8, 2015, Volume 113:
Some organochlorine pesticides and some chlorophenyl herbicides
Oct 5-13, 2015, Volume 114:
Rodent and processed meat
Monograph Working Group
Members
A. Blaz (USA) - Meeting Chair;
I. Enechi (Australia);
L. McCann (UK);
G.M. Calzavara (Canada);
G.M. Calzavara (Canada);
F. Oesch (Germany);
P. Vineis (Italy);
J. Ferlinz (France);
H. Kawanishi (Japan);
A. T. Mwanuzi (New Zealand);
T. Ruzsinszky (Hungary);
P. Eggerth (Ireland) (unable to attend)

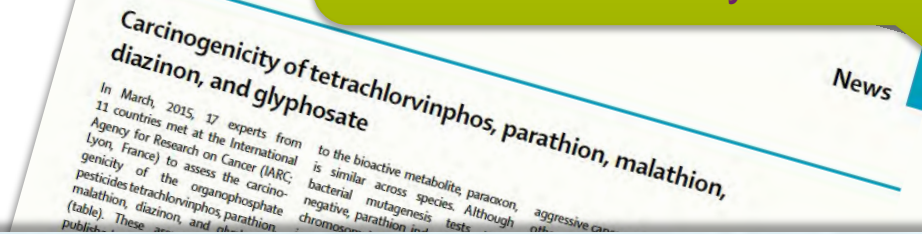
Pesticide	Activity (current status)	Evidence in humans (cancer sites)			Classification ^a
		Inadequate	Sufficient	Mechanistic evidence	
Tetrachlorvinphos	Insecticide (restricted in the EU and for most uses in the USA)	Inadequate	Sufficient	-	2B
Parathion	Insecticide (restricted in the USA and EU)	Inadequate	Sufficient	-	2B
Malathion	Insecticide (currently used; high production volume chemical)	Limited (non-Hodgkin lymphoma, prostate)	Sufficient	Genotoxicity, oxidative stress, inflammation, receptor-mediated effects, and cell proliferation or death	2A†
Diazinon	Insecticide (restricted in the USA and EU)	Limited (non-Hodgkin lymphoma, leukemia, lung)	Limited	Genotoxicity and oxidative stress	2A†
Glyphosate	Herbicide (currently used; highest global production volume herbicide)	Limited (non-Hodgkin lymphoma)	Sufficient	Genotoxicity and oxidative stress	2A†

EU-European Union. ^a See the International Agency for Research on Cancer (IARC) preamble for explanation of classification system (summarized in January 2006). † The 2A classification of diazinon was based on limited evidence of carcinogenicity in humans and experimental animals, and strong mechanistic evidence; for malathion and glyphosate, the mechanistic evidence provided independent support of the 2A classification based on evidence of carcinogenicity in humans and experimental animals.

Bestrijdingsmiddelen kankerverwekkend?



Hazard (Not Risk!) Characterization
 Class 2B: Possibly carcinogenic in humans
 Class 2A: Probably carcinogenic in humans



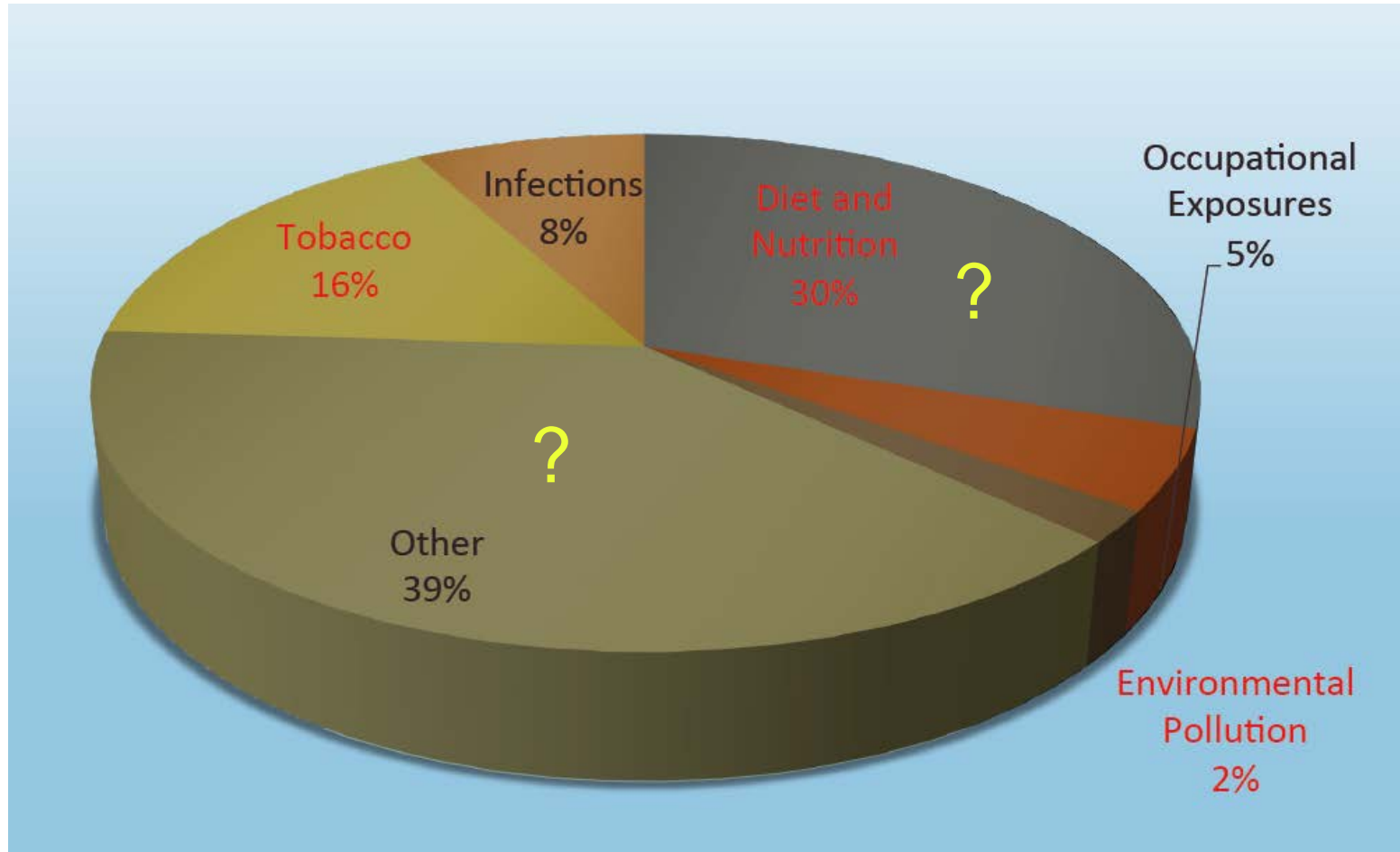
	Activity (current status)	Evidence in humans (cancer sites)	Evidence in animals	Mechanistic evidence	Classification*
Tetrachlorvinphos	Insecticide (restricted in the EU and for most uses in the USA)	Inadequate	Sufficient	..	2B
Parathion	Insecticide (restricted in the USA and EU)	Inadequate	Sufficient	..	2B
Malathion	Insecticide (currently used; high production volume chemical)	Limited (non-Hodgkin lymphoma, prostate)	Sufficient	Genotoxicity, oxidative stress, inflammation, receptor-mediated effects, and cell proliferation or death	2A†
Diazinon	Insecticide (restricted in the USA and EU)	Limited (non-Hodgkin lymphoma, leukaemia, lung)	Limited	Genotoxicity and oxidative stress	2A†
Glyphosate	Herbicide (currently used; highest global production volume herbicide)	Limited (non-Hodgkin lymphoma)	Sufficient	Genotoxicity and oxidative stress	2A†

EU=European Union. *See the International Agency for Research on Cancer (IARC) preamble for explanation of classification system (amended January, 2006). †The 2A classification of diazinon was based on limited evidence of carcinogenicity in humans and experimental animals, and strong mechanistic evidence; for malathion and glyphosate, the mechanistic evidence provided independent support of the 2A classification based on evidence of carcinogenicity in humans and experimental animals.

Table: IARC classification of some organophosphate pesticides

Causes for Cancer in Industrialized Countries (2006)

American Cancer Society and IARC-WHO



Pesticides [66]		Pesticides	Chem	Pesticides	Chem	Pesticides	Chemical structure
2,4-D (H) M(g/Mol) = 221 pKa = 2.73 logP: 2.81		Aldrin (I) M(g/Mol) = 364.9 pKa = n.a logP: 6.5		Bioallethrin (I) M(g/Mol) = 302.4 pKa = n.a logP: 4.68		Carbendazim (F) M(g/Mol) = 191.2 pKa = 4.2 logP: 1.48	
Acephate (I) M(g/Mol) = 183.2 pKa = 8.35 logP: -0.85		Atrazine (H) M(g/Mol) = 215.7 PKa = 4.14, 10.7 logP: -0.97		Bitertanol (F) M(g/Mol) = 337.4 pKa = n.a logP: 4.1		Carbofuran (I) M(g/Mol) = 151.1 pKa = n.a logP: 1.48	
Acetochlor (H) M(g/Mol) = 269.8 pKa = n.a logP: 4.14							

Int. J. Environ. Res. Public Health 2011, 8, 2265-2303; doi:10.3390/ijerph8062265

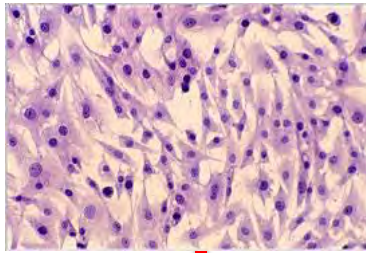
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International Journal of
Environmental Research and
Public Health
ISSN 1660-4601
www.mdpi.com/journal/ijerph

Review Effect of Endocrine Disruptor Pesticides: A Review

Wisssem Mnif^{1,2}, Aziza Ibn Hadj Hassine¹, Aicha Bouaziz¹, Aghleb Bartegi³, Olivier Thomas⁴
and Benoit Roig^{4,*}

Chloromethoxy (I) M(g/Mol) = 505.2 pKa = n.a logP: 4.6		Endrin (I) M(g/Mol) = 380.9 pKa = n.a logP: 3.2		Fenbuconazole (F) M(g/Mol) = 336.8 pKa = n.a logP: 3.79	
Cypermethrin (I) M(g/Mol) = 416.3 pKa = n.a logP: 5.3		Diazinon (I) M(g/Mol) = 327.4 pKa = 2.6 logP: 3.69		Fenitrothion (I) M(g/Mol) = 277.2 pKa = n.a	
Cyproconazole (F) M(g/Mol) = 291.8 pKa = n.a logP: 3.09		Dichlorvos (I) M(g/Mol) = 221 pKa = n.a logP: 1.9		Fenvalerate (I) M(g/Mol) = 419.9 pKa = n.a logP: 5.01	

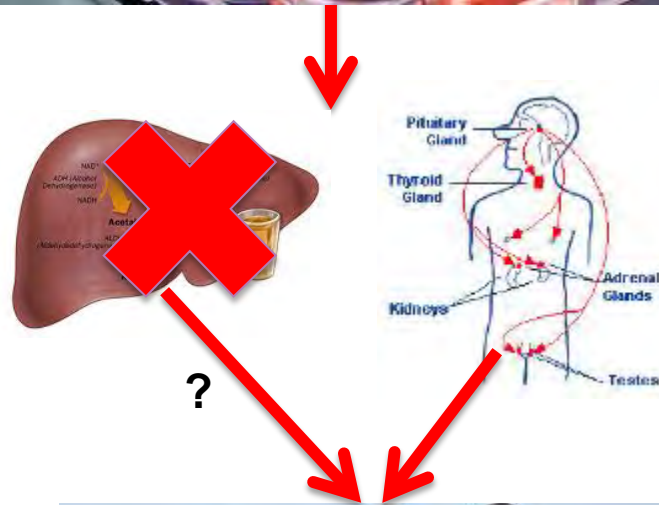
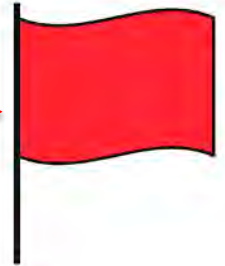
Nog geen 50% van genoemde bestrijdingsmiddelen die in het review artikel genoemd worden!



Genotoxiciteit



Endocriene
verstoring?



Risicoschatting





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Zwakke punten in
experimentele
(dier)modellen?



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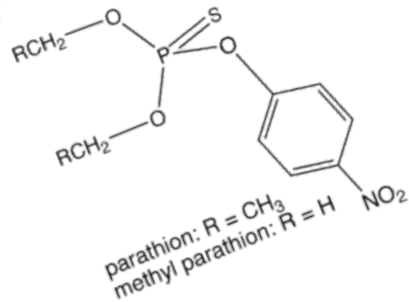
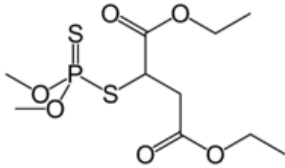
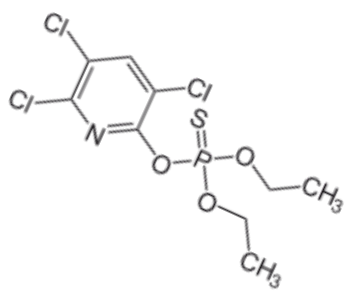
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Risico's van
blootstelling in de
vroege
levensfase



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Hersenen ontwikkeling en gedrag bij mens slecht te voorspellen met proefdieronderzoek





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Groeiende aandacht
en zorg voor socio-
economische
(gezondheids)
effecten



Meer aandacht voor gezondheid vs economische kosten samenleving

Universiteit Utrecht

HEAL. (2014). *Health costs in the European Union - how much is related to EDCs?*. <http://www.env-health.org/>: Health and Environment Alliance (HEAL).

Trasande, L., Zoeller, R. T., Hass, U., Kortenkamp, A., Grandjean, P., Myers, J. P., . . . Heindel, J. J. (2015). Estimating burden and disease costs of exposure to endocrine-disrupting chemicals in the European Union. *The Journal of Clinical Endocrinology and Metabolism*, 100(4), 1245-1255. doi:10.1210/jc.2014-4324 [doi]

Olsson, I., Holmer, M. L., Carlsson, M., Kjäll, K., Ramböll, A. P., Niemelä, H., . . . Olsson, M. (2014). The cost of inaction-A socioeconomic analysis of costs linked to effects of endocrine disrupting substances on male reproductive health.

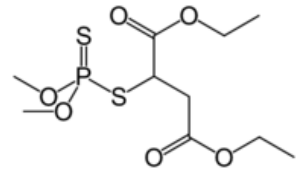
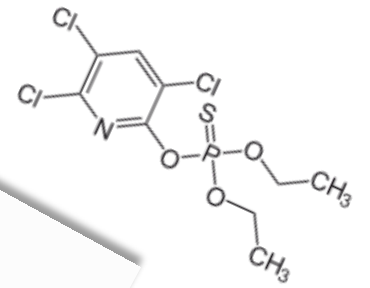
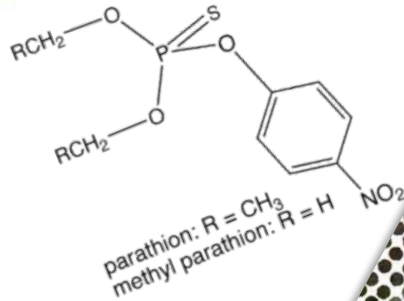
Rijk, I., & van den Berg, M. (June 10, 2015). *Putting a price on your exposed brain: A case-study towards prenatal exposure to polybrominated diphenyl ethers (PBDEs), organophosphate pesticides (OPs) and associated socioeconomic cost of IQ loss In The Netherlands*. The Netherlands: Institute for Risk Assessment Sciences (IRAS), Utrecht University.

Rijk, I., van Duursen M., & van den Berg, M. (2016) Health effects related to Endocrine Disrupting Chemicals and their socio-economic impact in the EU – An inventory, evaluation and way forward in cost estimates of EDC-related health effects (*in prep.*). The Netherlands: Institute for Risk Assessment Sciences (IRAS), Utrecht University.



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Institute for Risk Assessment Sciences

Putting a price on your exposed brain

A case-study towards prenatal exposure to polybrominated diphenyl ethers (PBDEs), organophosphate pesticides (OPs) and associated socio-economic cost of IQ loss in the Netherlands

Prof. dr. Ingrid Rijk, MSc
Prof. dr. Martin van den Berg

June 2015





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PEDIATRICS

OFFICIAL JOURNAL OF THE AMERICAN ACADEMY OF PEDIATRICS

Attention-Deficit/Hyperactivity Disorder and Urinary Metabolites of Organophosphate Pesticides

Maryse F. Bouchard, David C. Bellinger, Robert C. Wright and Marc G. Weisskopf
Pediatrics 2010;125:e1270; originally published online May 17, 2010;
DOI: 10.1542/peds.2009-3058



American Journal of Epidemiology
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Original Contribution

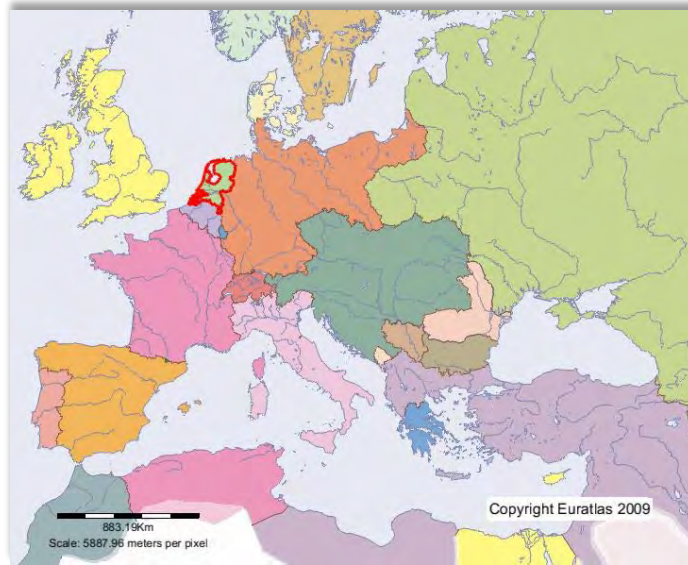
Prenatal Organophosphate Metabolite Performance on the Brazelton Neonatal Behavioral Assessment Scale in a Multiethnic Pregnancy Cohort

Stephanie M. Engel¹, Gertrud S. Berkowitz¹, Dana B. Barr², Susan L. Teitelbaum¹, Jodi Siskind¹,
Stefanie J. Meisel¹, James G. Wetmur^{3,4}, and Mary S. Wolff¹

Vol. 165, No. 12
DOI: 10.1093/aje/kwm029
Advance Access publication April 3, 2007



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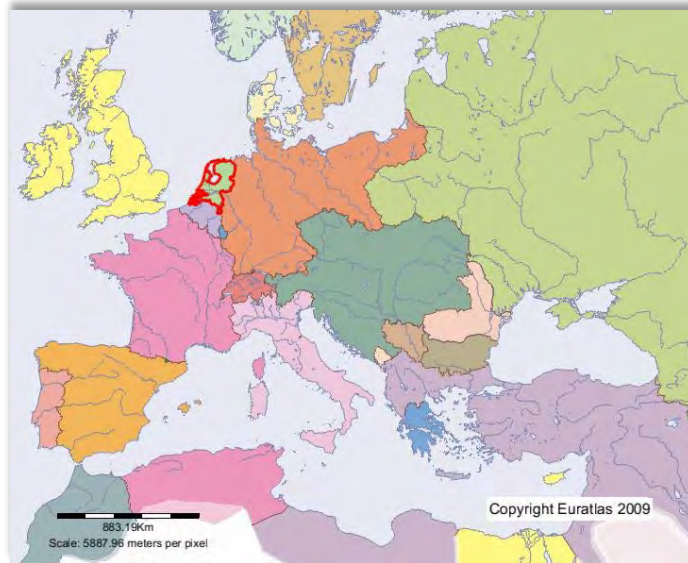
Concentraties OP-
pesticiden in NL
2000-2010



Table 6. Lost IQ points and associated socio-economic cost per year of OPs exposure for three exposure groups in the Netherlands. Calculations are based on the ERR from the CHAMACOS –study cohort (Bouchard et al., 2011)

Exposure group	Total DAP reference level (nmol/L)	IQ loss per newborn (95% C.I.)	Amount of newborns
P50-P75	272	-1.69 (-2.72; -0.66)	42 835
P75-P95	494	-3.14 (-5.04; -1.23)	34 268
>P95	1116	-5.12 (-8.23; -2.01)	8567
Total			

Ye, X., Pierik, F. H., Hauser, R., Duty, S., Angerer, J., Park, M. M., . . . Longnecker, M. P. (2008).. *Environmental Research*, 108(2), 260-267.



What are the socio-economic cost for the NL?

Table 6. Lost IQ points and associated socio-economic cost per year of OPs exposure for three exposure groups in the Netherlands. Calculations are based on the ERR from the CHAMACOS -study cohort (Bouchard et al., 2011)

Exposure group	Total DAP reference level (nmol/L)	IQ loss per newborn (95% C.I.)	Amount of newborns	IQ loss per exposure group	Socio-economic cost (95% C.I.) (million € / year of exposure)
P50-P75	272	-1.69 (-2.72; -0.66)	42 835	-116 070 (-172 713; -42 219)	-875 (-1407; -344)
P75-P95	494	-3.14 (-5.04; -1.23)	34 268	-107 466 (-172 713; -42 219)	-1303 (-2093; -512)
>P95	1116	-5.12 (-8.23; -2.01)	8567	-43 856 (-70 483; -17229)	-532 (-854; 209)
Total				-223 455 (-359 266; -87 821)	-2709 (-4354; -1064)

1 – 4 miljard € per jaar

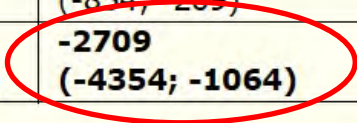


Table 9. Range of EDC-attributable cost per health effect and total EDC-attributable socio-economic cost the EU (in billion €). Outliers in cost estimates and their proposed alternatives are indicated in red

Source	Nordic Council			HEAL		Trasande			This /other reports			overall range		Proposal for best range	
	2%	20%	40%	Total (2%)	Total (5%)	low	base case	high	1% / low	2,5% / base case	10% / high	lowest	highest	lowest	highest
Reproductive tract and fertility															
Reduced female fertility				0,048 - 0,062	0,120 - 0,155										
Reduced male fertility	0,007	0,072	0,145			4,71	4,71	4,71				0,007	4,71	0,007	0,155
Cryptorchidism	0,018	0,181	0,363	0,018 - 0,026	0,045 - 0,065	0,117	0,130	0,250				0,018	0,363	0,018	0,363
Hypospadias	0,009	0,089	0,178									0,009	0,178	0,009	0,178
Endometriosis									0,775	1,94	7,75	0,775	7,75	0,775	7,75
Neurobehavioral diseases and disorders															
Autism spectrum disorders (ASD)				4,52	11,3	0,080	0,199	0,399				0,080	11,3	0,080	11,3
AD(H)D				0,014	0,035	2,62	4,14	4,93				0,014	4,93	2,62	4,93
IQ loss						42,2	133,4	183,6	32,0	84,3	136	42,2	183,6	42,2	183,6
Mental retardation						6,11	22,6	33,43				6,11	33,4	6,11	33,4
Neural tube defects									0,008	0,019	0,077	0,008	0,077	0,008	0,077
Hormone-related cancers															
Breast cancer				0,320	0,800							0,320	0,800	0,320	0,800
Prostate cancer				0,180	0,450							0,180	0,450	0,180	0,450
Testis (testicular germ cell)															
Metabolic syndroms, imm															
Obesity: Obe															
Obesity adult						15,6	15,6	15,6				1,62	17,2	1,62	17,2
Diabetes mellitus (type 2)				6,0	15,0	1,44	1,44	17,2				1,44	17,2	1,44	17,2
Increment death rate among men						7,96	7,96	7,96				7,96	7,96	0,8	0,8
Asthma									0,173	0,432	1,73	0,173	1,73	0,173	1,73
TOTAL (billion €)	0,059	0,591	1,185	12,7	31,6	44,7	192,6	270,4	NA	NA	NA	60,9	292,6	56,4	280,8
TOTAL (billion €) after correction	NA			NA		157 (90% C.I. 32 – 212)			NA			NA		NA	

Geschatte directe en indirecte kosten ED effects, including biociden, **50-300 miljard € op jaarbasis voor EU28**

Rijk, I., van Duursen M., & van den Berg, M. (2016) Health effects related to Endocrine Disrupting Chemicals and their socio-economic impact in the EU –An inventory, evaluation and way forward in cost estimates of EDC-related health effects (*in prep.*). The Netherlands: Institute for Risk Assessment Sciences(IRAS), Utrecht University.



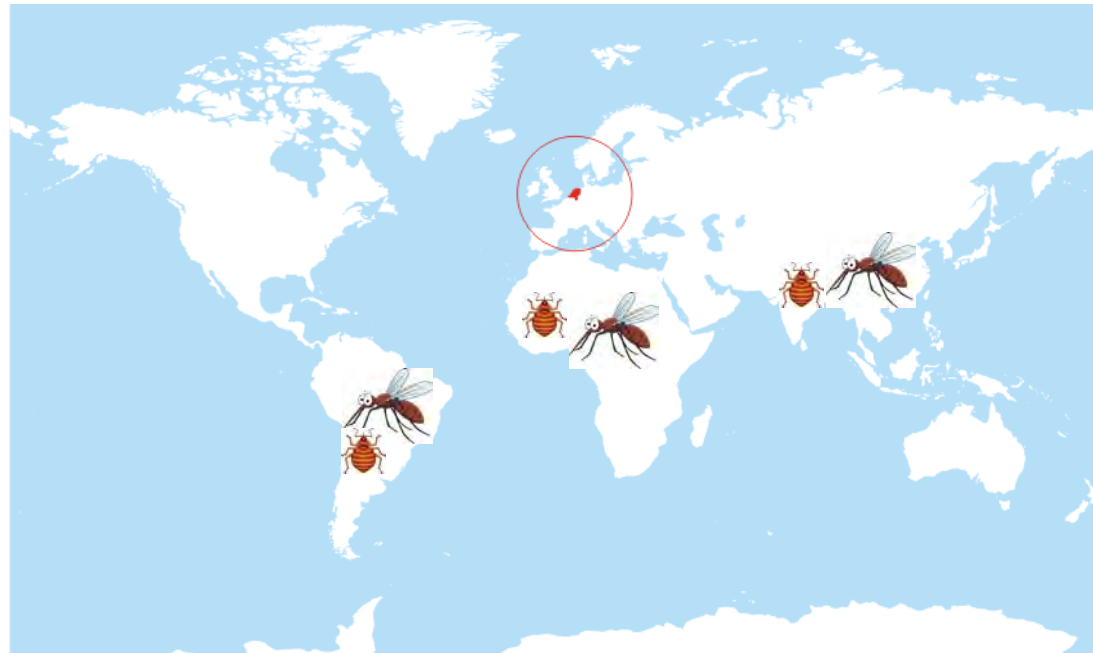
Socio-economische voordelen
wegen niet altijd op tegen de
nadelen vanuit
gezondheidsoogpunt





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“Risico-Voordeel”
analyses van biociden
in mondiaal perspectief



World Health
Organization





Mosquitoes:

- Chikungunya
- Dengue fever
- Rift Valley fever
- Yellow fever
- Zika
- Malaria
- Japanese encephalitis
- Lymphatic filariasis
- West Nile fever



Aquatic snails
Bilharziasis

Triatomine bugs
Chagas disease



Flies

Leishmaniasis
Sandfly fever
Sleeping sickness
River blindness



Fleas

Plague
Rickettsiosis



Ticks:

Haemorrhagic fever
Lyme disease
Relapsing fever
Spotted and Q fever)
Encephalitis
Tularaemia



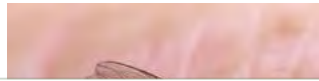


Mosquito

- Chikungunya
- Dengue
- Rift Valley fever
- Yellow fever
- Zika
- Malaria
- Japanese encephalitis
- Lymphatic filariasis
- West Nile virus



Flies



World Health Organization

Every year more than 1 billion cases and over 1 million deaths from vector-borne diseases such as malaria, dengue, schistosomiasis, human African trypanosomiasis, leishmaniasis, Chagas disease, yellow fever, Japanese encephalitis and onchocerciasis, globally.

Vector-borne diseases account for over 17% of all infectious diseases.



c fever
e
ver
Q fever)

Triatomine bugs

Chagas disease



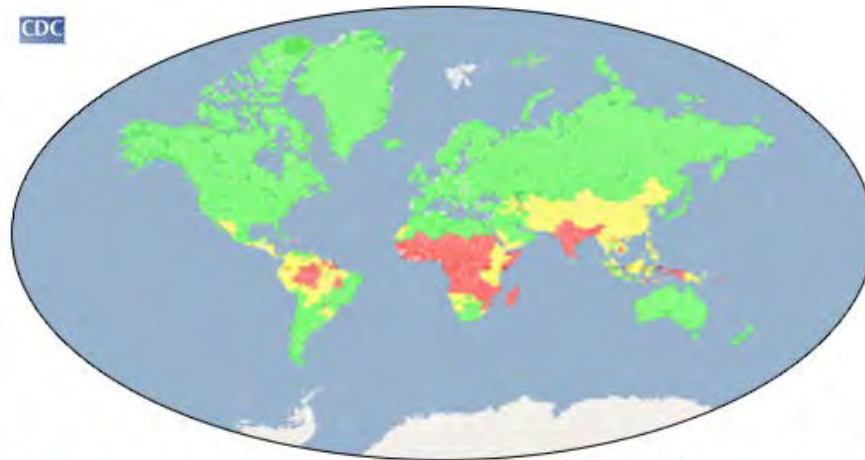



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CDC



 Malaria transmission occurs throughout

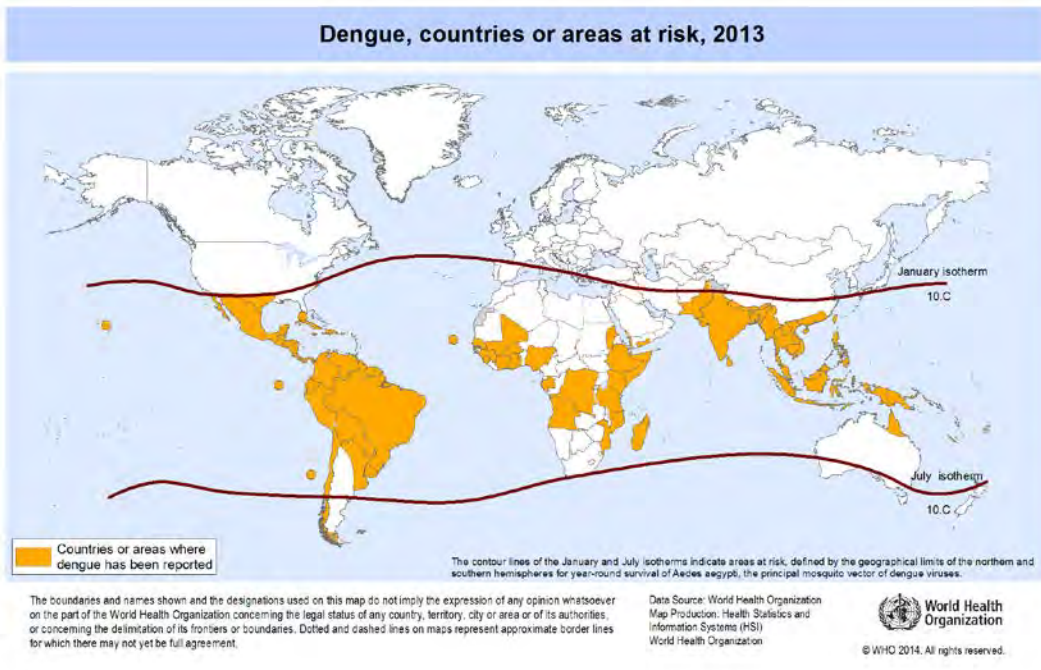
 Malaria transmission occurs in some parts

 Malaria transmission is not known to occur

Malaria

About 40% of the world's population at risk.
Up to 500 million cases occur every year
Up to 2.7 million deaths annually.

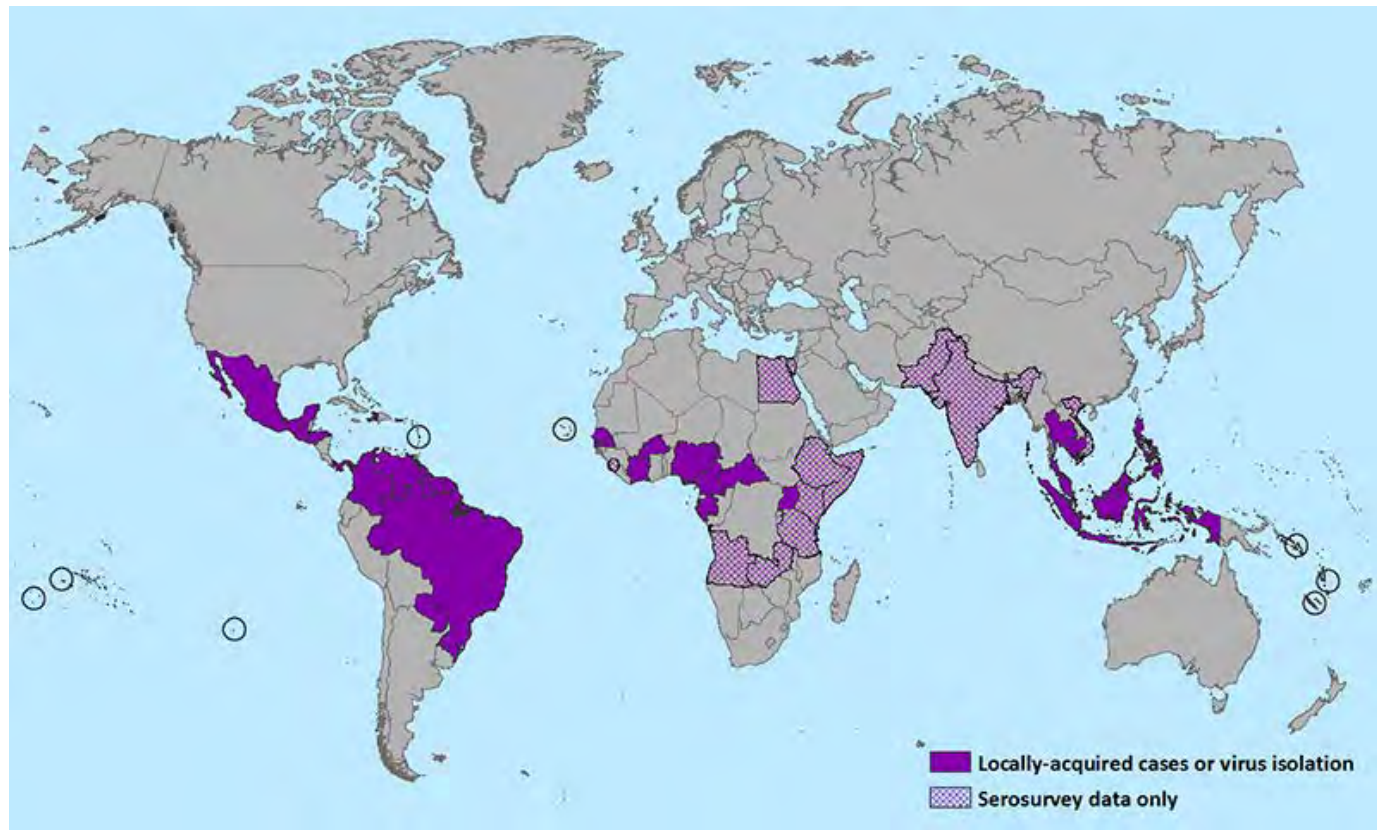




Dengue

2500 million people worldwide at risk of infection
 20 million cases a year in more than 100 countries.
 Tens of thousands serious dengue haemorrhagic fever.





Zika virus distribution

Symptomen vergelijkbaar met Dengue en Chikungunya

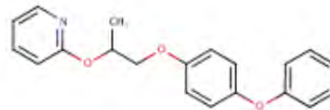
WHOPES-recommended compounds and formulations for control of mosquito larvae



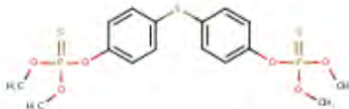
World Health Organization

Insecticide compounds and formulation(s) ¹	Class group ²	Dosage (active ingredient)		
		General (open water bodies)		Container-breeding (mg/L)
		(g/ha)	(mg/m ²)	
<i>Bacillus thuringiensis israelensis</i> , strain AM65-52, WG (3000 ITU/mg)	BL	125–750 ³	12.5–75 ³	1–5 ³
<i>Bacillus thuringiensis israelensis</i> , strain AM65-52, GR (200 ITU/mg)	BL	5,000–20,000 ³	500–2000 ³	-
Chlorpyrifos EC	OP	11–25	1.1–2.5	-
Diflubenzuron DT, GR, WP	BU	25–100	2.5–10	0.02–0.25
Novaluron EC	BU	10–100	1–10	0.01–0.05
Pyriproxyfen GR	JH	10–50	1–5	0.01
Fenthion EC	OP	22–112	2.2–11.2	-
Pirimiphos-methyl EC	OP	50–500	5–50	1
Temephos EC, GR	OP	56–112	5.6–11.2	1
Spinosad DT, EC, GR, SC	SP	20–500	2–50	0.1–0.5
Spinosad 83.3 monolayer DT	SP	250–500	25–50	-
Spinosad 25 extended release GR				
<i>Open bodies of water</i>	SP	250–400	25–40	-
<i>Control of Culex quinquefasciatus in open bodies of water with high organic matter</i>	SP	1000–1500	100–150	-

PYRIPROXYFEN
CASRN: 95737-68-1



TEMEPHOS
CASRN: 3383-96-8



Bacillus thuringiensis



Binnen Europa insecten als vector voor infectie-ziekten van zeer beperkt belang.
Hoe zit het dan met risico's van biociden op groente en fruit



Tabel 3: Controlegegevens per land van herkomst (op alfabetische volgorde)

2013-15

Land	aantal monsters	% boven wettelijke norm	middelen per monster
Argentinië	62	3,2	2,8
België	60	3,3	2,3
Brazilië	231	7,8	2,4
Chili	154	1,3	2,8
China	776	6,4	3,3
Colombia	76	11,8	2,0
Costa Rica	80	1,3	1,4
Dominicaanse Republiek	188	15,4	1,7
Duitsland	45	0,0	1,6
Egypte	455	10,3	3,4
Frankrijk	63	1,6	1,1
India	187	8,6	4,2
Israël	110	6,4	1,5
Italië	122	3,3	1,9
Kenya	1686	4,9	1,4
Marokko	201	3,5	2,0
Mexico	55	1,8	1,8
Nederland	1346	2,2	1,3
Onbekend	55	1,8	0,9
Peru	510	2,9	3,5
Spanje	727	2,5	2,1
Suriname	98	15,3	1,1
Thailand	139	17,3	1,4
Turkije	80	7,5	3,3
Verenigde Staten	48	2,1	1,6
Vietnam	100	30,0	2,4
Zuid-Afrika	353	3,7	2,2
Genoemde landen	7962	5,4	
totaal	8389	5,7	



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Nederlandse Voedsel- en
Warenautoriteit
Ministerie van Economische Zaken

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Biociden op groente en fruit : Een Risico?

- Meeste overschrijdingen van buiten de EU (ca 3%)
- Eenmalige overschrijding in voedsel minimaal risico
- Risico zit in chronische overschrijding
- Kwetsbaarste groepen zwangere vrouwen, kinderen
- Biologische producten geen meerwaarde voor algemene volksgezondheid (perceptie) vanuit biociden oogpunt
- Meerwaarde is minder biociden in milieu en daardoor minder impact op biodiversiteit



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Risico's biociden van uit **humaan** toxicologisch perspectief

- Moderne biociden onmisbaar voor bestrijding door insect-overdraagbare ziekten.
- “Risico-voordeel” principe zonder meer aan de voordeel kant bij insect-overdraagbare ziekten
- Risico's biociden op groente en fruit voor merendeel niet of minimaal, (wel voor veelvuldige blootstelling boven de norm gezondheidsrisico en kwetsbare groepen)
- De risico-perceptie hoger dan de feitelijke **Grootste onzekerheid!** in fruit over het algemeen
maar.....
- Huidige testmethoden risicos tijdens zwangerschap en eerste levensjaren vrijwel zeker te simpel en hebben maar beperkte voorspellende waarde

Risico's biociden van uit **eco**-toxicologisch perspectief

- Moderne biociden nog steeds impact op “non-target” organismen.
- Moderne biociden, in combinatie met monoculturen, kunnen een significante impact op biodiversiteit en ecosystemen hebben.
- In Nederland en EU landen de “luxe” om ons te bekommeren over eco-effecten. Grootste deel wereldbevolking heeft deze “luxe” niet door voedselschaarste, natuurrampen, klimaat etc.
- Rijke landen (w.o. NL) voortouw nemen in ontwikkeling ecologisch en ecotoxicologisch verantwoorde landbouw productie-methoden, die uiteindelijk ook toepasbaar zijn in andere delen van de wereld.





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Dank voor uw aandacht