

# Synopsis of published literatures on health effects of Endosulfan

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1) NERVOUS SYSTEM	
1.1	<p><b>Title:</b> Developmental exposure to pesticides zineb and/or endosulfan renders the nigrostriatal dopamine system more susceptible to these environmental chemicals later in life</p> <p><b>Authors and their affiliations:</b> Zhenquan Jia a, Hara P. Misra a,b,*</p> <p>a Department of Biomedical Sciences and Pathobiology, College of Veterinary Medicine, Virginia Polytechnic Institute and State University, Blacksburg, VA 24061, USA</p> <p>b Edward Via Virginia College of Osteopathic Medicine, Virginia Tech Corporate Research Center, 2265 Kraft Drive, Blacksburg, VA 24060, USA</p> <p><b>Published in:</b> NeuroToxicology 28 (2007) 727–735</p>
	<p><b>Major Findings:</b></p> <p>The study was to test the hypothesis that exposure to pesticides such as endosulfan and/or zineb during critical periods of postnatal development could result in neuronal dysfunction and enhance the impact of these pesticides during exposure as adults.</p> <p>The findings support our hypothesis that <b>exposure</b> to pesticides such as endosulfan and zineb during critical periods of postnatal development contributes to neurotransmitter changes upon re-challenge in adulthood.</p>
1.2	<p><b>Title:</b></p> <p><b>Major Findings:</b></p>

	<p><b>Endosulfan Poisoning and Chronic Brain Syndrome</b></p> <p><b>Authors and their affiliations:</b>  Dov R. Aleksandrowicz  Shalvata Psychiatric Center  Hod Hasharon, Israel</p> <p><b>Published in:</b>  <b>Arch. Toxicol. 43, 65-68 (1979)</b></p>	<p>This report describes a case of chronic brain syndrome following poisoning by <b>endosulfan</b>; it is believed to be <b>the first recorded case of such a complication due to that compound.</b></p>
1.3	<p><b>Title:</b>  <b>Effects of in utero and lactational exposure to endosulfan in prefrontal cortex of male rats</b></p> <p><b>Authors and their affiliations:</b>  Teresa Cabaleiro, Ana Caride,  Alejandro Romero, Anunciación Lafuente *  Laboratorio de Toxicología, Facultad de Ciencias, Universidad de Vigo, Campus de Orense, Las Lagunas, 32004 Orense, Spain</p> <p><b>Published in:</b>  <b>Toxicology Letters 176 (2008) 58–67</b></p>	<p><b>Major Findings:</b></p> <p>The possible <b>neurotoxic effects</b> of the organochlorine pesticide <b>endosulfan</b> have been <b>evaluated on male offspring rats exposed in utero and during lactation.</b></p> <p>The developing nervous system is proposed to be a potentially sensitive target for pesticide exposure (Tilson, 1998, 2000; Nakai and Satoh, 2002; Shafer and Meyer, 2005)</p> <p>Summarizing, <b>in utero and lactational treatment to endosulfan induces several alterations in content and metabolism of amino acids and biogenic amines in prefrontal cortex</b> in male rats during the development. These variations are age- and dose-</p>

		<p>dependent and <b>they could be related to several alterations in the important functions in which the prefrontal cortex is involved such as cognitive tasks, selective attention, short term working memory, response inhibition, behavioural flexibility, etc.</b></p>
1.4	<p><b>Title:</b>  <b>Effects of endosulfan on brain acetylcholinesterase activity in juvenile bluegill sunfish</b></p> <p><b>Authors and Affiliations:</b>  Hiran M. Dutta_ and Dane A. Arends  Department of Biological Sciences,  Kent State University, Kent, OH 44242,  USA</p> <p><b>Published in:</b> Environmental Research 91 (2003) 157–162</p>	<p><b>Major Points:</b></p> <p>The effects of endosulfan upon brain acetylcholinesterase (AChE) activity were measured in juvenile bluegill sunfish (<i>Lepomis macrochirus</i>).</p> <p><b>The duration of exposure was related to the reduction in the AChE activities which reflected the biotoxicity of endosulfan.</b></p> <p>The changes in the AChE activities will certainly affect the normal behavior of the juvenile bluegill which is <b>detrimental to their very existence in the natural habitat.</b></p>
1.5	<p><b>Title:</b>  <b>Selective involvement of basal ganglia and occipital cortex in a patient with acute endosulfan poisoning</b></p> <p><b>Authors and Affiliations:</b>  Department of Neurology, Sunjay Gandhi Postgraduate Institute of Medical Sciences, Lucknow, India</p> <p>Department of Radiology, Sunjay Gandhi Postgraduate Institute of Medical Sciences, Lucknow, India</p>	<p><b>Major Points:</b></p> <p>We report a teenage girl with acute endosulfan poisoning who developed psychosis, generalized tonic-clonic seizures, myoclonic jerks, cortical blindness and limb rigidity. Serial magnetic resonance imaging (MRI) <b>showed bilateral reversible lesions localized to caudate nucleus, putamen and occipital cortex;</b> internal capsule and thalamus were spared. This selective involvement may have a bearing on mechanisms underlying endosulfan toxicity.</p>

<p>Department of Neuroophthalmology, Sanjay Gandhi Postgraduate Institute of Medical Sciences, Lucknow, India</p> <p><b>Published in:</b> <b>Journal of Neurological Sciences 147</b> <b>(1997) 209-213</b></p>	
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<b>2) REPRODUCTIVE SYSTEM</b>		
2.1	<p><b>Title:</b> <b>A method to determine residue levels of persistent organochlorine pesticides in human milk from Indonesian women</b></p> <p><b>Authors and their affiliations:</b> Emma R. Burke a, Alexis J. Holden a,*, Ian C. Shaw b</p> <p>a Department of Environmental Management, University of Central Lancashire, Preston PR1 2HE, UK</p> <p>b Environmental Science &amp; Research Institute, 27 Creyke Rd, Christchurch, New Zealand</p> <p><b>Published in:</b> <b>Chemosphere 50 (2003) 529–535</b> 2002 Elsevier Science Ltd</p>	<p><b>Major Findings:</b></p> <p>A method has been developed for the analysis of organochlorine pesticides in human milk using solvent extraction, Florisil_ solid phase extraction clean-up and analysis by gas chromatography with an electron capture detector.</p> <p><b>There was no significant difference (at the 95% confidence level) in levels of pesticides between urban and rural areas</b></p>
2.2	<p><b>Title:</b> <b>Endosulfan sulphate interferes with reproduction, embryonic</b></p>	<p><b>Major Findings:</b></p> <p>The aim of this study was to evaluate the</p>

<p><b>development and sex differentiation in <i>Daphnia magna</i></b></p> <p><b>Authors and their affiliations:</b>  P. Palma a,c, V.L. Palmaa, R.M. Fernandes a, A.M.V.M. Soares b, I.R. Barbosa</p> <p>a Departamento de Cie^ncias do Ambiente, Escola Superior Agrá'ria de Beja, Beja 7800-295, Portugal</p> <p>b CESAM &amp; Departamento de Biologia da Universidade de Aveiro, Aveiro 3810-193, Portugal</p> <p>c Centro de Estudos Farmace^uticos, Faculdade de Farma'cia, Universidade de Coimbra, Rua do Norte, Coimbra 3000-295, Portugal</p> <p><b>Published in:</b>  <b>Ecotoxicology and Environmental Safety 72 (2009) 344– 350</b></p>	<p>possible effects promoted by endosulfan sulphate in changes on the life cycle, embryo development and sex differentiation of <i>Daphnia magna</i>.</p> <p><b>Endosulfan sulphate promoted a significant decrease of the offspring number in all concentrations.</b> Results showed a reduction of the size of females, together with a decrease in moulting frequency. Furthermore, <b>an increase in embryo deformities was observed at all concentrations tested.</b> Above a concentration of 91.7 mgL<sup>-1</sup> there was an increased production of males. The results suggest that <b>endosulfan sulphate interferes with the life cycle and sex determination</b> of the crustacean <i>D. magna</i>.</p>
<p>2.3 <b>Title:</b>  <b>Persistent Pesticides in Human Breast Milk and Cryptorchidism</b></p> <p><b>Authors and their affiliations:</b>  Ida N. Damgaard,<sup>1</sup> Niels E. Skakkebaek,<sup>1</sup> Jorma Toppari,<sup>2</sup> Helena E. Virtanen,<sup>2</sup> Heqing</p>	<p><b>Major Findings:</b></p> <p><b>Eight organochlorine pesticides</b> (including Endosulfan) were <b>measured</b> in all samples (medians; nanograms per gram lipid) for cases/controls.</p> <p>Statistical analysis showed that pesticide</p>

<p>Shen,<sup>3</sup> Karl-Werner Schramm,<sup>3</sup> Jørgen H. Petersen,<sup>1,4</sup> Tina K. Jensen,<sup>1</sup> Katharina M. Main</p> <p>1University Department of Growth and Reproduction, Copenhagen, Denmark;  2Departments of Physiology and Pediatrics, University of Turku, Turku, Finland;  3GSF-National Research Center for Environmental and Health, Institute for Ecological Chemistry, euerberg, Germany;  4Department of Biostatistics, University of Copenhagen, Denmark</p> <p><b>Published in:</b>  <b>Environ Health Perspect</b>  <b>114:1133–1138 (2006).</b></p>	<p>levels in breast milk were significantly higher in boys with cryptorchidism (<math>p = 0.032</math>).</p> <p>In conclusion, our study suggests an <b>association between congenital cryptorchidism and persistent organochlorine pesticides present in mothers' breast milk.</b></p> <p><b>Prenatal exposure to persistent organochlorine pesticides may adversely affect testicular descent in boys.</b></p>
<p>2.4 <b>Title:</b>  <b>Organochlorine pesticide residue levels and oxidative stress in preterm delivery cases</b></p> <p><b>Authors and their affiliations:</b>  Rahul Pathak<sup>1</sup>, Sanvidhan G Suke<sup>1</sup>, Tanzeel Ahmed<sup>1</sup>, Rafat S Ahmed<sup>1</sup>, AK Tripathi<sup>1</sup>, Kiran Guleria<sup>2</sup>, CS Sharma<sup>3</sup>, SD Makhijani<sup>3</sup> and BD Banerjee<sup>1</sup></p> <p>1 Environmental Biochemistry and Immunology laboratory, Department of Biochemistry,</p>	<p><b>Major Findings:</b></p> <p>The aim of the present study was to analyze the OCP residues in maternal and cord blood of women and assess the levels of different non-enzymatic oxidative stress markers as well as to establish correlation with OCP levels, if any.</p> <p>Levels of OCPs like Hexachlorocyclohexane (HCH), <b>endosulfan</b>, and DDT were analyzed by gas chromatography. Non-enzymatic oxidative stress was measured.</p> <p>In conclusion, our results suggest that</p>



<p>University College of Medical Sciences &amp; G.T.B. Hospital (University of Delhi), Delhi, India 2 Department of Obstetrics and Gynecology, University College of Medical Sciences &amp; G.T.B. Hospital (University of Delhi), Delhi, India</p> <p>3 Instrumentation and Bio-Labs, Central Pollution Control Board, Ministry of Environment and Forest, Delhi, India</p> <p><b>Published in:</b> <b>Human and Experimental Toxicology 29(5) 351–358</b></p>	<p>higher levels of some of the OCP (like <b>Endosulfan</b>) <b>may be associated with Preterm Delivery and increased oxidative stress.</b></p>
<p>2.5 <b>Title:</b> <b>Endosulfan and its metabolites in fertile women, placenta, cord blood, and human milk</b></p> <p><b>Authors and their affiliations:</b> Isabel Cerrilloa, Alicia Granadaa, Mari´a-Jose´ Lo´pez-Espinosaa, Begon˜a Olmosa, Margarita Jim´enez, Africa Can˜ob, Nicolas Oleaa,_, Mari´a Fa´tima Olea-Serranoc aLaboratory of Medical Investigations, Department of Radiology, School of Medicine, Hospital Clinico, University of Granada, 18071 Granada, Spain bGynecology &amp; Obstetric Service,</p>	<p><b>Major Findings:</b></p> <p>The present study <b>investigated the presence of endosulfan I, endosulfan II, and endosulfan metabolites in fatty and non-fatty tissues</b> and fluids from women of reproductive age and children in Southern Spain.</p> <p>The highest concentration of commercial endosulfan I and endosulfan II was found in adipose tissue, with a mean value (I+II) of 17.72 ng/g lipid, followed by human milk, with a mean value (I+II) of 11.38 ng/mL milk.</p> <p>This study suggests that <b>exposure from mother to child is a common event, both</b></p>

	<p>Hospital Clí'nico, Granada, Spain  Food Sciences Department,  University of Granada, Granada,  Spain</p> <p><b>Published in:</b>  <b>Environmental Research 98</b>  <b>(2005) 233–239</b></p>	<p><b>in utero and via breastfeeding</b>, due to the high frequency of exposure of women of reproductive age.</p>
2.6	<p><b>Title:</b>  <b>Effect of Endosulfan on Male Reproductive Development</b></p> <p><b>Authors and Affiliations:</b>  Habibullah Saiyed,<sup>1</sup> Aruna Dewan,<sup>1</sup> Vijay Bhatnagar,<sup>1</sup> Udyavar Shenoy,<sup>2</sup> Rathika Shenoy,<sup>2</sup> Hirehall Rajmohan,<sup>3</sup> Kumud Patel,<sup>1</sup> Rekha Kashyap,<sup>1</sup> Pradip Kulkarni,<sup>1</sup> Bagalur Rajan,<sup>3</sup> and Bhadabhai Lakkad<sup>1</sup></p> <p><sup>1</sup>National Institute of Occupational Health (Indian Council of Medical Research), Meghani Nagar, Ahmedabad, India;</p> <p><sup>2</sup>Department of Pediatrics, Kasturba Medical College, Mangalore, India; <sup>3</sup>Regional Occupational Health Research Centre, Bangalore, India</p> <p><b>Published in:</b>  <b>VOLUME 111   NUMBER 16  </b>  <b>December 2003 • Environmental Health Perspectives</b></p>	<p><b>Major Points:</b></p> <p>A study to examine the relationship between environmental endosulfan exposure and reproductive development in male children and adolescents.</p> <p>The study population was composed of 117 male schoolchildren (10–19 years of age) of a village situated at the foothills of cashew plantations.</p> <p>The study parameters included recording of clinical history, physical examination, sexual maturity rating (SMR) according to Tanner stages, and estimation of serum levels of testosterone, luteinizing hormone (LH), follicle-stimulating hormone, and endosulfan residues.</p> <p>Our study <b>results suggest that endosulfan exposure in male children may delay sexual maturity and interfere with sex hormone synthesis.</b></p>



<b>3) ENDOCRINE SYSTEM</b>		
3.1	<p><b>Title:</b>  <b>Effect of pesticides on estrogen receptor transactivation in vitro: A comparison of stable transfected MVLN and transient transfected MCF-7 cells</b></p> <p><b>Authors and their affiliations:</b>            Eva C. Bonefeld-Jorgensen *, Heidi T. Grunfeld, Irene M. Gjermansen</p> <p>Unit of Environmental Biotechnology, Department of Environmental and Occupational Medicine, University of Aarhus, Vennelyst Boulevard 6, DK-8000 Aarhus, Denmark</p> <p><b>Published in:</b>  <b>Molecular and Cellular Endocrinology 244 (2005) 20–30</b></p>	<p><b>Major Findings:</b></p> <p>The <b>estrogenic potential of four pesticides</b> (endosulfan, prochloraz, tolchlofos-methyl and propamocarb) was <b>compared</b> in parallel with 17_estradiol (E2) by reporter constructs in transient transfected MCF-7BUS and in stable transfected MVLN cells.</p> <p><b>In MVLN cells, endosulfan, prochloraz, tolchlofos-methyl and propamocarb caused cytotoxic responses at concentrations higher than 10, 25, 25 and 100 _M, respectively.</b></p>
3.2	<p><b>Title:</b>  <b>Xenoestrogen-Induced ERK-1 and ERK-2 Activation via Multiple Membrane-Initiated Signaling Pathways</b></p> <p><b>Authors and their affiliations:</b>            Bulayeva, Nataliya N;Watson, Cheryl S</p> <p>Department of Human Biological Chemistry and genetics, University of Texas Medical Branch, Galveston,</p>	<p><b>Major Findings:</b></p> <p>The objective of the study was to investigate the ability of some of the estrogen mimetics (belonging to major classes of environmental estrogens) to produce rapid activation of ERKs via various signaling pathways in the GH3/B6/F10 prolactinoma cell lines.</p> <p><b>Endosulfan was able to produce phosphorylation at almost all tested concentrations.</b></p> <p>Basically two patterns of stimulations were seen: Compounds active <b>in both</b></p>

	<p>Texas, USA</p> <p><b>Published in:</b>  <b>Environmental Health Perspectives; Nov 2004; 112, 15; ProQuest pg. 1481</b></p>	<p><b>subpicomolar and nanomolar ranges</b> (Endosulfan, nonylphenol and coumestrol) versus compounds active only in the nanomolecular range (DDE and dieldrin).</p>
<p>3.3</p>	<p><b>Title:</b>  <b>Effects of Currently Used Pesticides in Assays for Estrogenicity, Androgenicity, and Aromatase Activity in Vitro</b></p> <p><b>Authors and their affiliations:</b>  Helle Raun Andersen,* Anne Marie Vinggaard,† Thomas Høj Rasmussen,* Irene Marianne Gjermandsen,‡ and Eva Cecilie Bonefeld-Jørgensen‡</p> <p>*Environmental Medicine, Institute of Public Health, University of Southern Denmark, Odense, Winsløwparken 17, DK-5000 Odense C, Denmark;  †Institute of Food Safety and Toxicology, Division of Biochemical and Molecular Toxicology, Danish Veterinary and Food Administration, Mørkhøj Bygade 19, DK-2860 Søborg, Denmark; and  ‡Department of Environmental and Occupational Medicine, Unit of Environmental Biotechnology, University of Aarhus, Building 260, Vennelyst Boulevard 6, DK-8000 Aarhus C, Denmark</p>	<p><b>Major Findings:</b></p> <p><b>Twenty-four pesticides were tested for interactions with the estrogen receptor (ER) and the androgen receptor (AR) in transactivation assays.</b></p> <p><b>The two organochlor insecticides dieldrin and Endosulfan increased cell proliferation and ER transactivation gene response in MCF-7 cells significantly at 5 and 1 <math>\mu</math>M, respectively. Both compounds induced maximum response at 25 <math>\mu</math>M, whereas higher concentrations resulted in a decreased response due to cytotoxicity. In both assays, the response induced by endosulfan was higher than the response induced by dieldrin. Endosulfan potentiated the 17<math>\beta</math>-estradiol-induced proliferation when it was tested together with a concentration of 17<math>\beta</math>-estradiol causing a submaximum response. Dieldrin did not enhance the 17<math>\beta</math>-estradiol-induced response in any of the assays.</b></p>

	<p><b>Published in:</b>  <b>Toxicology and Applied Pharmacology 179, 1–12 (2002)</b></p>	
3.4	<p><b>Title:</b>  <b>Endosulfan effects on pituitary hormone and both nitrosative and oxidative stress in pubertal male rats</b></p> <p><b>Authors and Affiliations:</b>  A. Caride, A. Lafuente*, T. Cabaleiro  Laboratorio de Toxicología, Facultad de Ciencias, Universidad de Vigo, Campus de Orense, Las Lagunas, 32004 Orense, Spain</p> <p><b>Published in:</b>  <b>Toxicology Letters 197 (2010) 106–112</b></p>	<p><b>Major Points:</b>  The study was undertaken to investigate in pubertal male rats possible effects of endosulfan administered throughout lactation and gestation on: (a) pituitary gene expression of prolactin, luteinizing hormone (LH), growth hormone (GH) and thyroid stimulating hormone (TSH); and (b) circulating levels of these hormones.</p> <p>In pubertal male rat, prenatal and lactational exposure to endosulfan modifies expression and release of prolactin, LH, GH and TSH, and pituitary NOS1 and NOS2 mRNA levels, suggesting that nitrosative stress can be implicated in the endocrine toxicity of endosulfan at pituitary level.</p>
3.5	<p><b>Title:</b>  <b>Screening of some anti-progestin endocrine disruptors using a recombinant yeast based in vitro bioassay</b></p> <p><b>Authors and Affiliations:</b>  Shamba Chatterjee a,1, Vikas Kumar a,1, Chandrajeet B. Majumder b, Partha Roy a,*  a Molecular Endocrinology Laboratory, Department of Biotechnology, Indian Institute of Technology Roorkee, Roorkee 247 667, Uttarakhand, India</p> <p>b Fluid Particle Research Laboratory, Department of</p>	<p><b>Major Points:</b>  About 7 different chemicals (mostly pesticides or their metabolites) like DDT and its metabolites, nonylphenol, endosulfan were screened in this assay system for their role in transactivation and they were all found to be anti-progestative.</p>

	<p>Chemical Engineering, Indian Institute of Technology Roorkee, Roorkee 247 667, Uttarakhand, India</p> <p><b>Published in:</b> <b>Toxicology in Vitro 22 (2008) 788–798</b></p>	
3.6	<p><b>Title:</b> Circulating thyroid hormone levels and iodothyronine deiodinase activities in Nile tilapia (<i>Oreochromis niloticus</i>) following dietary exposure to Endosulfan and Aroclor</p> <p><b>Authors and Affiliations:</b> Ana Maria Coimbra<sup>a,b,*</sup>, Maria Armanda Reis-Henriques<sup>a,b</sup>, Veerle M. Darrasc</p> <p>a) ICBAS-Instituto de Cie<sup>^</sup>ncias Biome<sup>^</sup>dicas de Abel Salazar, Laborato<sup>^</sup>rio de Fisiologia Aplicada, Largo do Prof. Abel Salazar 2, 4099-003, Porto, Portugal</p> <p>b) CIIMAR-Centro Interdisciplinar de Investigac<sup>~</sup>o Marinha e Ambiental, Laborato<sup>^</sup>rio de Toxicologia Ambiental, Rua dos Bragas 289, 4050-123, Porto, Portugal</p> <p>c) Laboratory of Comparative Endocrinology, Zoological Institute K.U. Leuven, Naamsestraat 61, B-3000 Leuven, Belgium</p> <p><b>Published in:</b> <b>Comparative Biochemistry and Physiology, Part C 141 (2005) 8 – 14</b></p>	<p><b>Major Points:</b> We evaluated the effects of two organochlorinated environmental contaminants, Endosulfan and Aroclor 1254 on peripheral thyroid hormone metabolism and thyroid hormone plasma levels in Nile tilapia (<i>Oreochromis niloticus</i>).</p> <p>It is concluded that dietary exposure of tilapia to <b>Endosulfan</b> or Aroclor 1254 <b>can lead to changes in circulating thyroid hormone levels and/or in peripheral thyroid hormone metabolism.</b> The changes in hormone metabolism differ between tissues, eventually reflecting tissue-specific differences in adaptation.</p>

3.7	<p><b>Title:</b>  <b>Human Exposure to Endocrine-Disrupting Chemicals and Prenatal Risk Factors for Cryptorchidism and Hypospadias: A Nested Case-Control Study</b></p> <p><b>Authors and Affiliations:</b>  Mariana F. Fernandez,<sup>1</sup> Begoña Olmos,<sup>1</sup> Alicia Granada,<sup>1</sup> Maria José López-Espinosa,<sup>1</sup> José-Manuel Molina-Molina,<sup>1</sup> Juan Manuel Fernandez,<sup>2</sup> Milagros Cruz,<sup>3</sup> Fátima Olea-Serrano,<sup>4</sup> and Nicolás Olea<sup>1</sup></p> <p><sup>1</sup>Laboratory of Medical Investigations, <sup>2</sup>Department of Pediatrics, and <sup>3</sup>Department of Gynecology and Obstetrics, San Cecilio University Hospital, Granada, Spain; <sup>4</sup>Department of Nutrition, University of Granada, Granada, Spain</p> <p><b>Published in:</b>  <b>VOLUME 115   SUPPLEMENT 1   December 2007 • Environmental Health Perspectives</b></p>	<p><b>Major Points:</b></p> <p>In this study we aimed to determine whether the combined effect of environmental estrogens measured as total effective xenoestrogen burden (TEXB) is a risk factor for male urogenital malformations.</p> <p><b>We found an increased risk for male urogenital malformations related to the combined effect of environmental estrogens in placenta.</b></p>
3.8	<p><b>Title:</b>  <b>Endocrine disruptors and human puberty</b></p> <p><b>Authors and Affiliations:</b>  E. Den Hond and G. Schoeters</p> <p>Department of Toxicology, VITO (Flemish Institute for Technological Research), Mol, Belgium</p> <p><b>Published in:</b>  <b>international journal of andrology 29 (2006) 264–271</b></p>	<p><b>Major Points:</b></p> <p>Epidemiological research studying the effect of endocrine disruptors on the onset of puberty is summarized.</p> <p><b>In girls, earlier age at menarche</b> was reported after exposure to polychlorinated biphenyls (PCBs), polybrominated biphenyls (PBBs), persistent pesticides (DDT, endosulfan) and phthalate esters.</p> <p><b>In boys,</b> exposure to PCBs, PCDFs or the pesticide <b>endosulfan</b> was associated with <b>delayed puberty or decreased penile length.</b></p>

3.9	<p><b>Title:</b>  <b>Oxidative stress and loss of cortisol secretion in adrenocortical cells of rainbow trout (<i>Oncorhynchus</i>) exposed in vitro to endosulfan, an organochlorine pesticide</b></p> <p><b>Authors and Affiliations:</b>  J. Dorval, V.S. Leblond, A. Hontela *</p> <p>De'partement des Sciences Biologiques, Universite' du Que'bec a` Montre'al, TOXEN Research Centre, C.P. 8888, succ. Centre-ville, Montre'al, Que', Canada H3C 3P8</p> <p><b>Published in:</b>  <b>Aquatic Toxicology 63 (2003) 229_/241</b></p>	<p><b>Major Points:</b>  The effects of endosulfan on cortisol secretion, cell viability, antioxidants and lipid peroxidation were investigated.</p> <p>The in vitro study identified endosulfan as a chemical inducing a loss of secretory responses in teleost adrenocortical steroidogenic cells and alterations in the activity of enzymes known to be involved in oxidative stress pathways.</p> <p>Moreover, the significant increase in lipid hydroperoxides levels provided further evidence for endosulfan-induced oxidative stress.</p>
3.10	<p><b>Title:</b>  <b>Effect of in vitro estrogenic pesticides on human oestrogen receptor _ and _ mRNA levels</b></p> <p><b>Authors and Affiliations:</b>  H.T. Grünfeld, E.C. Bonefeld-Jorgensen*</p> <p><i>Unit of Environmental Biotechnology, Department of Environmental and Occupational Medicine, University of Aarhus, Vennelyst Boulevard 6, DK-8000 Aarhus, Denmark</i></p> <p><b>Published in:</b>  <b>Toxicology Letters 151 (2004) 467–480</b></p>	<p><b>Major Points:</b>  Nine widely distributed pesticides were recently demonstrated to possess potential estrogenic properties in oestrogen receptor (ER) transactivation and/or E-screen assays.</p> <p>This study demonstrated that organochlor and organo-phosphorous pesticides possess the ability to interfere with the ER_ and ER_ mRNA steady state levels.</p>



<b>4) IMMUNE SYSTEM</b>		
4.1	<p><b>Title:</b></p> <p><b>Determination of the immunotoxic potential of pesticides on functional activity of sheep leukocytes in vitro</b></p> <p><b>Authors and Affiliations:</b>            Juraj Pistl *, Nata'lia Kovalkovic'ova', Vanda Holovska', Jaroslav Lega'th, Ivan Mikula</p> <p>Department of Microbiology and Immunology, University of Veterinary Medicine, 041 81 Kos'ice, Slovakia</p> <p><b>Published in:</b> Toxicology 188 (2003) 73_/81</p>	<p><b>Major Points:</b></p> <p>The effect of eight pesticides with different chemical structure (atrazine, bentazone, chloride-zone, dichlofluanid, <b>endosulfan</b>, MCPA, simazine, triallate) on sheep peripheral blood phagocytes and lymphocytes was examined.</p> <p>Three of the pesticides tested <b>suppressed both, the metabolic activity of phagocytes and mitogenic activation of lymphocytes</b> (dichlofluanid, endosulfan and simazine).</p>
4.2	<p><b>Title:</b></p> <p><b>Toxic effects of endosulfan on blood lymphocyte subsets in adult rats</b></p> <p><b>Authors and Affiliations:</b>            A. Lafuente, T. Cabaleiro, A. Caride, A. Romero  <i>Laboratorio de Toxicolog'ia, Facultad de Ciencias, Universidad de Vigo, Campus de Orense, Las Lagunas s/n, 32004-Orense, Spain</i></p> <p><b>Published in:</b>  <b>Toxicology Letters 164S (2006) S1-S324</b></p>	<p><b>Major Points:</b></p> <p>The objective was to evaluate possible immune alterations induced by endosulfan exposure during pregnancy and lactation in both male and female adult rats.</p> <p>The results suggest that <b>endosulfan exposure</b> during pregnancy and lactation could <b>modify humoral immunity</b> in offspring adulthood, and that <b>females</b> are <b>more susceptible</b> to this pesticide because their cellular immunity was also altered.</p>

**5) GENOTOXICITY, CYTOTOXICITY AND TERATOGENICITY**

<p>5.1</p>	<p><b>Title :</b></p> <p><b>Comet assay in phytoplankton as biomarker of genotoxic effects of environmental pollution</b></p> <p><b>Authors and their affiliations:</b></p> <p>Akcha a,* , G. Arzul a, S. Rousseau a, M. Bardouil b</p> <p>aIFREMER, Laboratoire d'Ecotoxicologie, Rue de l'Île d'Yeu, 44311 Nantes Cedex 03, France</p> <p>b IFREMER, Laboratoire Phycotoxines, Rue de l'Île d'Yeu, 44311 Nantes Cedex 03, France</p> <p><b>Published in:</b></p> <p><b>Marine Environmental Research 66 (2008) pg: 59–61</b></p> <p>journal homepage:  <a href="http://www.elsevier.com/locate/marenvrev">www.elsevier.com/locate/marenvrev</a>          _ 2008 Elsevier Ltd.</p>	<p><b>Major Findings:</b></p> <p><b>Genotoxicity was observed from 1 lg/L of endosulfan and was not concentration dependent.</b></p> <p><b>Endosulfan exposure resulted in DNA strand breaks from the concentration of 1 lg/L (p &lt; 0.05), but effect did not appear to be concentration dependent</b></p>
<p>5.2</p>	<p><b>Title :</b></p> <p><b>Endosulfan Decreases Cell Growth and Apoptosis in Human HaCaT Keratinocytes:Partial ROS-Dependent ERK1/2 Mechanism</b></p> <p><b>Authors and their affiliations:</b></p> <p>SEBASTIEN ANThERIEU,1* NATHALIE LEDIRAC,1 ANNE-PASCALE LUZY,2 PHILIPPE LENORMAND,3 JEAN-CLAUDE CARON,2</p>	<p><b>Major Findings:</b></p> <p>This study was designed to assess the mitogenic, apoptogenic, and genotoxic effects of endosulfan on the HaCaT cell line.</p> <p><b>Endosulfan</b> has been shown to generate transient reactive oxygen species (ROS), and blocking this oxidative stress by</p>

	<p>AND ROGER RAHMANI*</p> <p>1 INRA, UMR 1112 “Re’ponses des Organismes aux Stress Environnementaux”, Equipe de Toxicologie Cellulaire, Mole’culaire et Ge’nomique, Sophia-Antipolis, France</p> <p>2 Galderma R&amp;D, Les Templiers, Biot, France</p> <p>3 Institute of Signaling, Developmental Biology and Cancer Research, CNRS-UMR 6543, Centre Antoine Lacassagne, Nice, France</p> <p><b>Published in:</b></p> <p><b>J. Cell. Physiol. 213: 177–186, 2007.</b></p>	<p>N-acetyl cysteine (NAC) strongly prevented both persistent nuclear ERK1/2 phosphorylation and cell growth decrease. Additional experiments demonstrated that unchanged endosulfan rather than its metabolites has <b>mutagenic</b> effects and <b>increased DNA strand breaks</b> in human keratinocytes.</p> <p>In conclusion, the results show that human keratinocytes are <b>more affected by a chronic</b> than an acute <b>exposure</b> to endosulfan.</p> <p>On the whole, our findings demonstrate that endosulfan <b>disrupts epidermal homeostasis by two concomitant independent mechanisms:</b> a ROS-dependent mechanism responsible for DNA damage and decrease in cell proliferation, and a ROS-independent inhibition of apoptosis that could contribute to mutant cell survival and therefore have possible carcinogenic effects.</p>
5.3	<p><b>Title :</b></p> <p><b>Analysis of Endosulfan and Its Metabolites in Human Serum Using Gas Chromatography–Tandem Mass Spectrometry</b></p>	<p><b>Major Findings:</b></p> <p>A new analytical method combining solid-phase extraction and gas chromatography–tandem mass spectrometry used to determine the insecticide endosulfan as well as its</p>

	<p><b>Authors and their affiliations:</b></p> <p>F.J. Arrebola<sup>1</sup>, J.L. Martínez Vidal<sup>1,*</sup>, and A. Fernández-Gutiérrez<sup>2</sup></p> <p><sup>1</sup> Department of Analytical Chemistry, University of Almería, 04071 Almería, Spain</p> <p><sup>2</sup> Department of Analytical Chemistry, University of Granada, 18071 Granada, Spain</p> <p><b>Published in:</b></p> <p><b>Journal of Chromatographic Science, Vol. 39, May 2001</b></p>	<p>metabolites endosulfan-ether, -lactone, and -sulfate in human serum.</p> <p>The <b>mean levels of the total endosulfan</b> found in the farmers were <b>significantly higher</b> than the mean level present in the non-occupationally exposed people group.</p> <p>The results obtained revealed that the pesticide could be detected practically intact in human blood, the similarity of the results obtained from the serum samples of the two studied groups and the differences found when the <b>urine samples</b> were analyzed seemed to <b>indicate</b> that urine analysis offers information about a <b>recent exposition</b> of the individual and serum analysis shows that information can be attributed to less recent expositions.</p>
5.4	<p><b>Title :</b></p> <p><b>Oxidative stress responses in different organs of <i>Jenynsia multidentata</i> exposed to Endosulfan</b></p> <p><b>Authors and their affiliations:</b></p> <p>M.L. Ballesterosa, D.A.Wunderlin<sup>b</sup>, M.A.Bistonía,</p> <p>a Universidad Nacional de Córdoba, Facultad de Ciencias Exactas y Físicas</p>	<p><b>Major Findings:</b></p> <p>Our present results demonstrate that exposure to a sublethal concentration of <b>Endosulfan</b> results in oxidative stress in several organs of <i>J. multidentata</i>.</p> <p>The <b>brain</b> was the <b>most sensitive organ to oxidative damage</b>.</p>

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5.5	<p><b>Title :</b></p> <p><b>DNA Damage and Mutagenicity Induced by Endosulfan and Its Metabolites</b></p> <p><b>Authors and their affiliations:</b></p> <p>Mahima Bajpayee,<sup>1</sup> Alok Kumar Pandey,<sup>1</sup> Sabina Zaidi,<sup>2</sup> Javed Musarrat,<sup>2</sup> Devendra Parmar,<sup>1</sup> Neeraj Mathur,<sup>3</sup> Prahlad Kishore Seth,<sup>1</sup> and Alok Dhawan<sup>1*</sup></p> <p>1) Developmental Toxicology Division, Industrial Toxicology Research Centre, Lucknow-226001, India</p> <p>2) Department of Microbiology, Faculty of Agricultural Sciences, Aligarh Muslim University, Aligarh-202002, India</p> <p>3) Epidemiology Section, Industrial Toxicology Research Centre, Lucknow-226001, India</p> <p><b>Published in:</b>  <b>Environmental and Molecular Mutagenesis 47:682–692 (2006)</b> VVC 2006 Wiley-Liss, Inc.</p>	<p><b>Major Findings:</b></p> <p>Endosulfan and metabolites may interact with DNA directly or through the production of ROS.</p> <p>In conclusion, the present study is the first to compare endosulfan with its isomers and metabolites for their ability to induce mutations in a conventional bacterial test system and to induce DNA damage in mammalian cells. Endosulfan, its isomers, and metabolites were positive in both the bacterial and mammalian cell systems</p>

5.6	<p><b>Title :</b></p> <p><b>Selective induction of the CYP3A family by endosulfan and DNA-adduct formation in different hepatic and hepatoma cells</b></p> <p><b>Authors and their affiliations:</b></p> <p>M. Dubois a, A. Pfohl-Leszkowicz ', I. De Waziers ', P. Kremers S*</p> <p>a Universit P de Litge. Lahorcrtoire de Chimie M:dicale, Institut de Pathologir, B35, Centre Hospitalier Unir~ersitaue. B-4000 Surf Tilman, Belgium</p> <p>h E.N.S.A.T.. Lahorrtrtoire de Toxicologic et SCcuritr' Alimentaire. 145 Al. de Muret. 31076 Toulouse. France</p> <p>' Uniti de Rerherche de Biochirnie Pharmacologicue et MPtabolique (INSERM U751. CHU Necker-Erzfant Malades. 156 rue de Vougirard. 75730 Paris Cedex 15. France</p> <p><b>Published in:</b></p> <p><b>Environmental Toxicology and Pharmacology 1 (1996) 249-256</b></p>	<p><b>Major Findings:</b></p> <p>This report clearly shows the <b>genotoxicity</b> of endosulfan. This chlorinated insecticide induces the formation of abundant DNA adducts in both fetal rat hepatocytes and Hep G2 cells.</p> <p>Endosulfan strongly correlated to the high induction of CYP3A gene expression.</p> <p>Endosulfan has a <b>major impact on the human liver</b> via induction of CYP3A4 and CYP3A7.</p>
5.7	<p><b>Title:</b></p> <p><b>Exposure to mixtures of endosulfan and zineb induces apoptotic and necrotic cell death in SH-SY5Y neuroblastoma cells, in vitro</b></p> <p><b>Authors and their affiliations:</b></p> <p>Zhenquan Jia<sup>1</sup> and Hara P. Misra<sup>1,2,*</sup></p> <p><sup>1</sup> Department of Biomedical Sciences and Pathobiology, College of Veterinary</p>	<p><b>Major Findings:</b></p> <p>A number of epidemiological studies have demonstrated a strong association between the incidence of Parkinson's disease and pesticide exposure</p> <p>In conclusion, our results help to better characterize pesticide-induced neuroblastoma cell death. <b>Both zineb and</b></p>



	<p>Medicine, Virginia Polytechnic Institute and State University, Blacksburg, VA 24061, USA</p> <p>2 Edward Via Virginia College of Osteopathic Medicine, Virginia Tech Corporate Research Center, 2265 Kraft Drive, Blacksburg, VA 24060, USA</p> <p><b>Published in:</b></p> <p><b>JOURNAL OF APPLIED TOXICOLOGY</b></p> <p><b>J. Appl. Toxicol. 2007; 27: 434–446</b></p> <p><b>Published online 16 February 2007 in Wiley InterScience</b></p> <p><b>(www.interscience.wiley.com) DOI: 10.1002/jat.1218</b></p>	<p><b>endosulfan were found to induce cytotoxicity in SH-SY5Y cells via both apoptotic and necrotic pathways.</b></p> <p>Together, with evidence provided in this study, it appears likely that endosulfan and zineb alone or in combination play a role in the cell death in an in vitro model system that may, at least in part, be involved in the <b>degeneration of dopaminergic neurons and may be relevant for the pathogenesis of some of these neurodegenerative diseases.</b></p>
5.8	<p><b>Title:</b></p> <p><b>Advantages of Human Hepatocyte-Derived Transformants Expressing a Series of Human Cytochrome P450 Isoforms for Genotoxicity Examination</b></p> <p><b>Authors and their affiliations:</b></p> <p>Tsuneo Hashizume,* Sumie Yoshitomi,† Satoru Asahi,* Rieko Uematsu,* Shigeo Matsumura,* Fumio Chatani,* and Hiroaki Oda‡,1</p> <p>*Development Research Center and †Discovery Research Center, Pharmaceutical Research Division, Takeda Pharmaceutical Company Limited, 17-85</p>	<p><b>Major Findings:</b></p> <p>Endosulfan was found to induce MN through the CYP3A4-mediated pathway</p> <p>Endosulfan sulfate was the <b>genotoxic</b> metabolite to induce MN ; this metabolite was considered to be produced in the HepG2 transformant expressing human CYP3A4 by the treatment with b-endosulfan.</p>

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5.9	<p><b>Title:</b></p> <p><b>Citrinin and endosulfan induced teratogenic effects in Wistar rats</b></p> <p><b>Authors and their affiliations:</b></p> <p>Nittin D. Singh,† Anil K. Sharma,* Prabhaker Dwivedi, Rajendra D. Patil and Manoj Kumar</p> <p>Division of Pathology, Indian Veterinary Research Institute, Izatnagar- 243122, India</p> <p><b>Published in:</b></p> <p><b>JOURNAL OF APPLIED TOXICOLOGY</b></p> <p><b>J. Appl. Toxicol. 2007; 27: 143–151</b></p> <p><b>Published online 22 December 2006 in Wiley InterScience (www.interscience.wiley.com) DOI: 10.1002/jat.1185</b></p> <p>2006 John Wiley &amp; Sons, Ltd</p>	<p><b>Major Findings:</b></p> <p>The present investigation was conducted to evaluate the teratogenic potential of citrinin (CIT) and endosulfan either alone or in combination in pregnant rats during gestational days 6–20.</p> <p><b>In the present study, endosulfan treatment induced</b></p> <p>fetal gross anomalies The internal hydrocephalus, cerebellar hypoplasia, microphthalmia, contracted and notched kidneys, multilobulated liver, dilated renal pelvis, incomplete ossification of skull bones, rib anomalies and sacral and caudal vertebrae agenesis were the important fetal malformations</p> <p>The dose of endosulfan (1 mg kg<sup>-1</sup> body weight) employed in the present study</p>

		<p>is lower than the dose established by the World Health Organization (WHO) for reproductive toxicity assessment in rats (WHO, 1984).</p> <p>Because of widespread use or contamination of CIT and endosulfan in the environment and its occurrence in the animal and <b>human</b> food (Anderson, 1995) and the presence of residues in the blood and milk samples of females (Sancewicz-Pach et al., 1997; Sanghi et al., 2003), <b>these might pose a potential risk to developing fetuses during pre-natal and post-natal life</b></p>
5.10	<p><b>Title:</b></p> <p><b>Human health risk assessment of endosulfan. I: Toxicology and hazard identification</b></p> <p><b>Authors and their affiliations:</b></p> <p>Marilyn H. Silva *, Sheryl L. Beauvais</p> <p>Department of Pesticide Regulation, California Environmental Protection Agency, Sacramento, CA 95812, USA</p> <p><b>Published in:</b></p> <p><b>Regulatory Toxicology and Pharmacology</b> <b>56 (2010) 4–17</b></p> <p>0273-2300/\$ - see front matter Published by Elsevier Inc. doi:10.1016/j.yrtph.2009.08.013</p>	<p><b>Major Findings:</b></p> <p><b>A primary risk assessment concern for endosulfan is that effects resulting from exposure during critical developmental stages (in utero, or to infants and children) will result in endocrine disruption and subsequent neurotoxicity, developmental or reproductive adverse effects that are irreversible.</b></p>

5.11	<p><b>Title:</b></p> <p><b>Genotoxicity evaluation of the insecticide endosulfan in the wetland macrophyte <i>Bidens laevis</i> L</b></p> <p><b>Authors and their affiliations:</b></p> <p>De'borá J. Pe'rez a,b,c, Mirta L. Menone b,c,* , Elsa L. Camadro a,c, Vi'ctor J. Moreno b</p> <p>a) Laboratorio de Gene'tica, Estacio'n Experimental Agropecuaria Balcarce INTA e Facultad de Ciencias Agrarias e UNMdP,CC 276, 7620 Balcarce, Argentina</p> <p>b) Laboratorio de Ecotoxicolog'ia, Departamento de Ciencias Marinas, Facultad de Ciencias Exactas y Naturales UNMdP, Funes 3350 (7600) Mar del Plata, Argentina</p> <p>c) Consejo Nacional de Investigaciones Cient'ficas y Te'cnicas (CONICET), Rivadavia 1917, 1033, Buenos Aires, Argentina</p> <p><b>Published in:</b></p> <p><b>Environmental Pollution 153 (2008) 695e698</b></p> <p>_ 2007 Elsevier Ltd. All rights reserved.</p>	<p><b>Major Findings:</b></p> <p>The higher proportion of laggards and vagrant chromosomes observed at 5 mg/L would indicate that <b>endosulfan interacts with the spindle interrupting normal chromosome migration.</b></p> <p>Endosulfan resulted genotoxic to <i>B. laevis</i>, a species of potential value for bioassays and in situ monitoring of environmental contamination by pesticides.</p>
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5.12	<p><b>Title:</b></p> <p><b>Genotoxicity evaluation of acute doses of endosulfan to freshwater teleost <i>Channa punctatus</i> (Bloch) by alkaline single-cell gel electrophoresis</b></p> <p><b>Authors and their affiliations:</b></p> <p>Sanjay Pandey, N.S. Nagpure, Ravindra Kumar, Shilpi Sharma, Satish K. Srivastava, Mahendra S. Verma</p> <p>National Bureau of Fish Genetics Resources, Indian council of Agriculture Research, Lucknow 226002, Uttar Pradesh, India</p> <p><b>Published in:</b></p> <p><b>Ecotoxicology and Environmental Safety 65 (2006) 56–61</b></p> <p>2005 Elsevier Inc.</p>	<p><b>Major Findings:</b></p> <p>The Indian freshwater air-breathing teleost fish <i>Channa punctatus</i> (Bloch) was exposed to acute concentrations of the organochlorine pesticide <b>Endosulfan</b>.</p> <p>The presence of <b>DNA strand breakage</b> in exposed specimens indicated the <b>genotoxic potential</b> of endosulfan.</p>
5.13	<p><b>Title:</b></p> <p><b>Genotoxic Effects of <math>\alpha</math>-Endosulfan and <math>\beta</math>-Endosulfan on Human HepG2 Cells</b></p> <p><b>Authors and their affiliations:</b></p> <p>Yuquan Lu,<sup>1</sup> Kanehisa Morimoto,<sup>1</sup> Tatsuya Takeshita,<sup>1</sup> Toru Takeuchi,<sup>1</sup> and Takeshi Saito<sup>2</sup></p>	<p><b>Major Findings:</b></p> <p>In this study, we examined the <b>genotoxicity</b> of endosulfan in vitro with a HepG2 cell line.</p> <p>After treating HepG2 cells with <math>\alpha</math>- or <math>\beta</math>-endosulfan for 1 hr, <b>DNA strand breaks were significantly induced</b> by <math>\alpha</math>-endosulfan at</p>

	<p>1Department of Social and Environmental Medicine, Osaka University Graduate School of Medicine, Suita, Osaka, Japan;</p> <p>2Laboratory of Environmental Biology, Department of Preventive Medicine, Hokkaido University School of Medicine, Sapporo, Japan</p> <p><b>Published in:</b></p> <p>Environmental Health Perspectives * VOLUME 108 1 NUMBER 6 1 June 2000</p>	<p>concentrations from <math>2 \times 10^{-4}</math> M to <math>1 \times 10^{-3}</math> M, and by <math>\gamma</math>-endosulfan at <math>1 \times 10^{-3}</math> M.</p> <p>The results of this study suggest that both endosulfan is <b>genotoxic</b> to HepG2 cells.</p>
5.14	<p><b>Title:</b></p> <p><b>Micronucleus induction in erythrocytes of the <i>Hyla pulchella</i> tadpoles (Amphibia: Hylidae) exposed to insecticide endosulfan</b></p> <p><b>Authors and their affiliations:</b></p> <p>Rafael C. Lajmanovich a,*, Mariana Cabagna b, Paola M. Peltzer a, Gabriela A. Stringhini c, Andr�es M. Attademo c</p> <p>a National Council for Scientific and Technical Research (CONICET), Faculty of Biochemistry and Biological Sciences, FBCB-UNL, Pje. El Pozo s/n (3000), Santa Fe, Argentina</p> <p>b Cathedra of Normal Morphology, Faculty of Biochemistry and Biological Sciences, FBCB-UNL,</p>	<p><b>Major Findings:</b></p> <p>Results obtained here <b>showed a genotoxic effect of the endosulfan on erythrocytes</b> of <i>H. puchella</i>.</p> <p>It important noticed that commercial formulations of endosulfan utilized, contains many <b>'inert' ingredients</b> that can increase the toxicity of the product when compared to the technical-grade material. Furthermore, 'inert' ingredients used in formulated pesticide</p>



	<p>Pje. El Pozo s/n (3000), Santa Fe, Argentina</p> <p>c High School of Health, Faculty of Biochemistry and Biological Sciences, EES-FBCB-UNL (3001), Santa Fe, Argentina</p> <p><b>Published in:</b></p> <p><b>Mutation Research 587 (2005) 67–72</b></p> <p>2005 Elsevier B.V. All rights reserved.</p> <p>doi:10.1016/j.mrgentox.2005.08.001</p>	<p>products are usually not disclosed and not included in most of the testing required in order to register these pesticides. Moreover, although ‘inert’ ingredients have no pesticide activity, they may be biologically active and sometimes the most toxic component of a pesticide formulation.</p>
5.15	<p><b>Title:</b></p> <p><b>Effects of endosulfan on B cells of Langerhans islets in rat pancreas</b></p> <p><b>Authors and their affiliations:</b></p> <p>Yusuf Kalender a,*, Suna Kalender b, Meltem Uzunhisarcikli a,</p> <p>Ay.se Ogutcu a, Fatma Açikgoz a, Dilek Durak c</p> <p>a Biology Department, Faculty of Arts and Science, Gazi University, 06500 Ankara, Turkey</p> <p>b Biology Department, Faculty of Education, Gazi University, 06500 Ankara, Turkey</p> <p>c Biology Department, Yozgat Faculty of Arts and Science, Erciyes University, Yozgat, Turkey</p>	<p><b>Major Findings:</b></p> <p><b>Chronic endosulfan administration for 2 months led to degenerative changes of various degrees of the pancreatic islets as well as the exocrine acini.</b></p> <p>Insufficiency of insulin hormone causes the increase of blood glucose level at the end of the 5th and 6th weeks, blood glucose level significantly increased. This shows that B cells couldn’t secrete adequate insulin. Biochemical studies show that <b>endosulfan affects integral proteins and receptors of cell membrane.</b></p>

	<p><b>Published in:</b></p> <p>Toxicology 200 (2004) 205–211</p>	<p>In this study <b>endosulfan</b> caused <b>vacuoles and swelling of mitochondria</b>. Blood glucose levels and ultrastructural changes in this study show that <b>endosulfan affects B cells</b> of pancreas in rats even if it is <b>under the LD50 dose level</b>. Endosulfan not only has toxic effects on mammalian and other animals but it causes pollution as well. Therefore, microbial insecticides which are effective on target organisms and don't cause pollution should be used instead of endosulfan.</p>
5.16	<p><b>Title:</b></p> <p><b>Effect of pesticides on estrogen receptor transactivation in vitro: A comparison of stable transfected MVLN and transient transfected MCF-7 cells</b></p> <p><b>Authors and their affiliations:</b></p> <p>Eva C. Bonefeld-Jorgensen *, Heidi T. Grunfeld, Irene M. Gjermansen</p> <p>Unit of Environmental Biotechnology, Department of Environmental and Occupational Medicine, University of Aarhus, Vennelyst Boulevard 6, DK-8000 Aarhus, Denmark</p> <p><b>Published in:</b></p>	<p><b>Major Findings:</b></p> <p>The <b>estrogenic potential of four pesticides</b> (endosulfan, prochloraz, tolchlofos-methyl and propamocarb) was <b>compared</b> in parallel with 17<sub>β</sub>-estradiol (E2) by reporter constructs in transient transfected MCF-7BUS and in stable transfected MVLN cells.</p> <p><b>In MVLN cells, endosulfan, prochloraz, tolchlofos-methyl and propamocarb</b> caused <b>cytotoxic responses</b> at <b>concentrations higher than 10,</b></p>

	<p><b>Molecular and Cellular Endocrinology 244 (2005) 20–30</b></p>	<p>25, 25 and 100 <math>\mu</math>M, respectively.</p>
<p>5.17</p>	<p><b>Title:</b></p> <p><b>Reactive oxygen species in in vitro pesticide-induced neuronal cell (SH-SY5Y) cytotoxicity: Role of NF<math>\kappa</math>B and caspase-3</b></p> <p><b>Authors and their affiliations:</b></p> <p>(SH-SY5Y) cytotoxicity: Role of NF<math>\kappa</math>B and caspase-3</p> <p>Zhenquan Jia, Hara P. Misra ©</p> <p>Edward Via Virginia College of Osteopathic Medicine, Virginia Tech Corporate Research Center, 2265 Kraft Drive, Blacksburg, VA 24060, USA</p> <p><b>Published in:</b></p> <p><b>Free Radical Biology &amp; Medicine 42 (2007) 288–298</b></p>	<p><b>Major Findings:</b></p> <p>Our laboratory has demonstrated that mice exposed to endosulfan (an organochlorine cyclodiene pesticide) and zineb (zinc ethylene bisdithiocarbamate (EBDC) fungicide) as juveniles and re-exposed at 8 months of age showed loss of dopamine (DA) in striatum. Mixtures of these pesticides also caused significantly increased levels of alphasynuclein, a major component of Lewy bodies and a hallmark of neurodegenerative diseases such as PD and Alzheimer's disease (AD).</p> <p>In summary, the results of the present study demonstrate that neuronal cells (SH-SY5Y) exposed to endosulfan and zineb individually or in a mixture increase the production of</p>

		<p>hydrogen peroxide as well as superoxide anion, decrease SOD GPX, and CAT enzyme levels and increase lipid peroxide levels.</p> <p>This <b>holds tremendous implication</b> for the derivation of risk assessment guidelines for human exposure to pesticides.</p>
5.18	<p><b>Title:</b></p> <p><b>Organochlorine pesticide residue levels and oxidative stress in preterm delivery cases</b></p> <p><b>Authors and their affiliations:</b></p> <p>Rahul Pathak<sup>1</sup>, Sanvidhan G Suke<sup>1</sup>, Tanzeel Ahmed<sup>1</sup>, Rafat S Ahmed<sup>1</sup>, AK Tripathi<sup>1</sup>, Kiran Guleria<sup>2</sup>, CS Sharma<sup>3</sup>, SD Makhijani<sup>3</sup> and BD Banerjee<sup>1</sup></p> <p>1) Environmental Biochemistry and Immunology laboratory, Department of Biochemistry, University College of Medical Sciences &amp; G.T.B. Hospital (University of Delhi), Delhi, India</p> <p>2) Department of Obstetrics and Gynecology, University College of Medical Sciences &amp; G.T.B. Hospital (University of Delhi), Delhi, India</p> <p>3) Instrumentation and Bio-Labs, Central Pollution Control Board, Ministry of</p>	<p><b>Major Findings:</b></p> <p>The aim of the present study was to analyze the OCP residues in maternal and cord blood of women and assess the levels of different non-enzymatic oxidative stress markers as well as to establish correlation with OCP levels, if any.</p> <p>Levels of OCPs like Hexachlorocyclohexane (HCH), <b>endosulfan</b>, and DDT were analyzed by gas chromatography. Non-enzymatic oxidative stress was measured.</p> <p>In conclusion, our results suggest that higher levels of some of the OCP (like <b>Endosulfan</b>) <b>may be associated with Preterm Delivery and increased oxidative stress.</b></p>

	<p>Environment and Forest, Delhi, India</p> <p><b>Published in:</b></p> <p><b>Human and Experimental Toxicology 29(5)</b> 351–358</p>	
5.19	<p><b>Title:</b></p> <p><b>HUMAN RED BLOOD CELL MEMBRANE DAMAGE BY ENDOSULFAN</b></p> <p><b>Authors and Affiliations:</b></p> <p>CLIVE SUNIL DANIEL, SARITA AGARWAL and SHYAM S. AGARWAL</p> <p><i>ICMR Centre for Advanced Research in Genetics, Genetics Unit, Department of Medicine, K.G. Medical College, Lucknow (India)</i></p> <p><b>Published in:</b></p> <p><b>Toxicology Letters, 32 (1986) 113- 118</b></p>	<p><b>Major Points:</b></p> <p>Endosulfan's in vitro toxicity on human red blood cell membrane was studied by staining with a fluorochrome dye, merocyanine-540 (MC-540) and Scanning Electron Microscopy (SEM).</p> <p>At a <b>concentration of 0.001 pg/ml (1 ppb) endosulfan was found to damage human red cell membranes</b> as demonstrated by fluorescence of 30-50% of red cells on staining with MC-540.</p> <p>At concentration of 1 pg/ml (1 ppm) the cells were markedly damaged.</p>

6) CARCINOGENCITY		
6.1	<p><b>Title:</b></p> <p>Carcinogenic potential of endosulfan and its metabolites based on a quantum chemical model</p> <p><b>Authors and their affiliations:</b></p> <p>C.N.G. Bedor <sup>a</sup>, R.J.L. Morais <sup>a</sup>, L.S. Cavalcanti <sup>a</sup>, J.V. Ferreira <sup>b</sup>, A.C. Pavão</p> <p>Universidade Federal do Vale do São Francisco, Av. José de Sá Maniçoba, S/N, Centro, 56304-205, Petrolina, PE, Brazil</p> <p><sup>b</sup> Instituto Federal de Alagoas, Rua Mizael Domingues, 75, Poço, 57020-600, Maceió, AL, Brazil</p> <p><sup>c</sup> Universidade Federal de Pernambuco, Av. Prof. Moraes Rego, 1235, Cidade Universitária, 50670-901, Recife, PE, Brazil</p> <p><b>Published in:</b></p> <p><b>Science of the Total Environment 408 (2010) 6281–6284</b></p> <p>2010 Elsevier B.V. All rights reserved.</p>	<p><b>Major Findings:</b></p> <p>.The <b>aim</b> of the present study was <b>to investigate the carcinogenic potential of endosulfan</b> and its metabolites through electronic parameters that characterize the action of carcinogens, the <b>findings of the present study indicate that the use of this pesticide represents a risk to the health of the general population, especially rural workers.</b></p> <p>The data from the present <b>study indicate that endosulfan and all its metabolites have carcinogenic potential</b>, based on the computational quantum chemical model. This places human health at risk, which is especially true for agriculture workers who are most exposed to this compound and therefore the most vulnerable.</p>
6.2	<p><b>Title:</b></p> <p><b>Breast cancer risk and the combined effect of environmental estrogens</b></p> <p><b>Authors and their affiliations:</b></p> <p>Jesu<sup>±</sup> s M. Ibarluzea<sup>1</sup>, Mariana F. Fern<sup>±</sup>ndez<sup>2</sup>, Loreto Santa-Marina<sup>1</sup>, Maria F. Olea-Serrano<sup>2</sup>, Ana M. Rivas<sup>2</sup>, Juan J.</p>	<p><b>Major Findings:</b></p> <p>The <b>estrogenicity</b> of adipose tissue extracts due to bioaccumulated <b>xenoestrogens</b> was associated with a <b>higher risk of breast cancer</b> in the leaner women, especially in the</p>



	<p>Aurrekoetxea1, Jose<sup>±</sup> Expo<sup>±</sup> sito3, Miguel Lorenzo4, Pablo Torne<sup>±</sup> 5, Mercedes Villalobos6, Vicente Pedraza6, Annie J. Sasco7 &amp; Nicolas Olea2,*</p> <p>1Department of Health Guipuzkoa, San Sebastia'n, Basque Country, Spain;</p> <p>2Laboratory of Medical Investigations, Hospital Cli'nico University of Granada, Granada, Spain;</p> <p>3Department of Oncology, Virgen de las Nieves University Hospital,m Granada, Spain;</p> <p>4Department of Surgery, Torrecardenas Hospital, Almeri'a, Spain;</p> <p><b>Published in:</b></p> <p><b>Cancer Causes and Control 15: 591–600, 2004. 591 _ 2004 Kluwer Academic Publishers. Printed in the Netherlands.</b></p>	<p>postmenopausal leaner group.</p> <p>Complex interactions between chemicals, endogenous or exogenous hormones and their natural ligands and receptors <b>may alter the internal homeostasis of the estrogenic environment of mammary tissue, leading to malignant transformation and cancer.</b></p>
6.3	<p><b>Title:</b></p> <p><b>Modulation of aromatase activity and mRNA by various selected pesticides in the human choriocarcinoma JEG-3 cell line</b></p> <p><b>Authors and their affiliations:</b></p> <p>Nathalie Laville a, Patrick Balaguer b, Francois Brion a, Nathalie Hinfray a, Claude Casellas c, Jean-Marc Porcher a, S'elim A'it-A'issa a,*</p> <p>a INERIS, Ecotoxicological Risk Assessment Unit, BP 2, F-60550 Verneuil-</p>	<p><b>Major Findings:</b></p> <p>Some pesticides may induce aromatase activity through transcriptional activation of the CYP19 gene</p> <p>When considering the physiologic consequences of the modulating effects on aromatase by pesticides, it is</p> <p>important to relate the present findings to the levels of</p> <p>human exposure</p>

	<p>en-Halatte, France</p> <p>b INSERM, U540, Molecular and Cellular Endocrinology of Cancers, F-34090 Montpellier, France</p> <p>c CNRS UMR 5569, University of Montpellier 1, BP 14 491, F-34093 Montpellier Cedex 5, France</p> <p><b>Published in:</b></p> <p><b>Toxicology 228 (2006) 98–108</b></p> <p>© 2006 Elsevier Ireland Ltd. All rights reserved.</p>	<p>We found out new <b>inducers of aromatase activity</b>, namely aldrin, chlordane, cypermethrine, methylparathion, <b>endosulfan</b>, methoxychlor, oxadiazon and metolachlor.</p> <p>The present study strengthens the view that <b>these chemicals</b> may contribute to the occurrence of <b>reproductive adverse effects in both human and aquatic organisms.</b></p>
6.4	<p><b>Title:</b></p> <p><b>Relation of Prediagnostic Serum Estrogen and Androgen Levels to Breast Cancer Risk.</b></p> <p><b>Authors and their affiliations:</b></p> <p>CJoanne F. Dorgan, Christopher Longcope, Hugh E. Stephenson, Jr., Roni T. Falk, Rosetta Miller, Charlene Franz, Lisa Kahle, William S. Campbell, Joseph A. Tangrea, and Arthur Schatzkin</p> <p>Division of Cancer Prevention and Control, National Cancer Institute, Bethesda, Maryland 20892-7326</p> <p>Departments of Obstetrics and Gynecology and Medicine. University of Massachusetts</p>	<p><b>Major Findings:</b></p> <p>Results support the hypothesis that prediagnostic serum estrogens and <b>androgens are related to the subsequent diagnosis of breast cancer in postmenopausal woman</b></p>

	<p>Medical School, Worcester, Massachusetts 01655</p> <p><b>Published in:</b></p> <p><b>Cancer epidemiology biomarkers and prevention</b></p> <p><b>Cancer Epidemiol Biomarkers Prev 1996;5:533-539. Published online July 1, 1996.</b></p>	
6.5	<p><b>Title:</b></p> <p><b>The insecticide endosulfan and its two stereoisomers promote the growth of altered hepatic foci in rats</b></p> <p><b>Authors and their affiliations:</b></p> <p><b>Ronny Fransson-Steen<sup>1,2,3</sup>, Sten Flodström<sup>1</sup> and Lars WMRngard<sup>1</sup></b></p> <p><b><sup>1</sup>institute of Environmental Medicine, Karolinska Institutet, Box 60208, S-104 01 Stockholm and <sup>2</sup>Department of Toxicology, Karolinska Institute*, Box 60400, S-104 01 Stockholm, Sweden</b></p> <p><b>Published in:</b></p> <p><b>Carcinogenesis vol.13 no. 12 pp.2299-2303, 1992</b></p>	<p><b>Major Findings:</b></p> <p>Endosulfan was studied for its ability to act as a tumour promoter in a two-stage, altered hepatic foci bioassay in male Sprague-Dawley rats.</p> <p>The results show that endosulfan and its two stereoisomers promote the development of altered hepatic foci, suggesting that endosulfan is a tumour-promoting agent acting by clonal expansion of initiated cells.</p> <p>Our results show that the insecticide endosulfan is a potential liver tumour promoter in a similar manner to structurally-related chlorinated cyclodienes.</p>
6.6	<p><b>Title:</b></p> <p><b>Effects of Pesticides on the Ratio of 160c/2- hydroxyestrone: A Biologic Marker of Breast Cancer Risk</b></p> <p><b>Authors and their affiliations:</b></p>	<p><b>Major Findings:</b></p> <p>Xenobiotic estrogens are external compounds with estrogenic activity that may thereby affect the risk of breast cancer. <b>This paper describes a mechanism by which</b></p>

	<p>H. Leon Bradlow,<sup>1</sup> Devra Lee Davis,<sup>2</sup> Gong Lin,<sup>1</sup> Daniel Sepkovic,<sup>1</sup> and Raj Tiwari</p> <p><sup>1</sup>Strang-Cornell Cancer Research Laboratory, New York, New York; <sup>2</sup>office of the Assistant Secretary for Health, Department of Health and Human Services, Washington, DC</p> <p><b>Published in:</b></p> <p><b>Environ Health Perspect 103(Suppl 7):147-150 (1995)</b></p>	<p><b>xenoestrogens may affect the development of breast cancer.</b></p> <p><b>These pesticides significantly increase the ratio of 16<math>\alpha</math>-OHE1/2-OHE, metabolites to values comparable to or greater than those observed after DMBA (the known rodent carcinogen 7,12-dimethylbenzyl anthracene (DMBA)). The ratio of 16<math>\alpha</math>-OHE1/2-OHE1 may provide a marker for the risk of breast cancer.</b></p>
6.7	<p><b>Title:</b></p> <p><b>Serum Sex Hormone Levels After Menopause and Subsequent Breast Cancer</b></p> <p><b>Authors and their affiliations:</b></p> <p>F. Berrino, P. Muti, A. Micheli, V. Krogh, G. Secreto, Istituto Nazionale Tumori, Milan, Italy;</p> <p>G. Bolelli, Istituto di Clinica Ostetrica e Ginecologica "P Sfameni," Universita di Bologna, Italy;</p> <p>R. Sciajno, Istituto Nazionale Tumori, Milan, and Istituto di Clinica Ostetrica e Ginecologica "P Sfameni," Universita di Bologna;</p> <p>P. Pisani, Unit of Descriptive Epidemiology, International Agency for Research on Cancer, Lyon, France;</p>	<p><b>Major Findings:</b></p> <p>In conclusion, this study has produced two important findings. First, in our population, <b>high levels of steroid sex hormones were found to be powerful predictors of breast cancer.</b> Second, although the association between breast cancer risk and testosterone levels was already known from case-control studies these results prospectively confirm this association.</p>

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6.8	<p><b>Title:</b></p> <p><b>Estrogen Receptor Activation via Activation Function 2 Predicts Agonism of Xenoestrogens in Normal and Neoplastic Cells of the Uterine Myometrium</b></p> <p><b>Authors and Affiliations:</b></p> <p><b>Deborah S. Hunter, Leslie C. Hodges, Peter M. Vonier, Robin Fuchs-Young, Marco M. Gottardis, and Cheryl L. Walker<sup>2</sup></b></p> <p><b>The University of Texas M. D. Anderson Cancer Center, Science Park Research Division, Smithville, Texas 78957 [D. S. H., L. C. H., R. F-Y., C. L. W.];</b></p> <p><b>The Tulane-Xavier Center for Bioenvironmental Research, Tulane University, New Orleans, Louisiana 70112 [P. M. V.]; and</b></p> <p><b>Bristol Myers Squibb, Endocrine Oncology Pharmaceutical Research Institute,</b></p>	<p><b>Major Findings:</b></p> <p>This report describes an in vitro/in vivo system for identifying the effects of Estrogen receptor ligands in the myometrium and elucidating their mechanism of action.</p> <p>The results suggest that some exogenous Estrogens (like endosulfan) may mimic the effects of endogenous estrogens on uterine leiomyoma and may contribute to a complex hormonal milieu that impacts both normal and neoplastic myometrium.</p>

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