MONITORING POPS IN PACIFIC REGION: A CASE STUDY FOR OCDD CONTAMINATION IN FIJI ISLANDS

 $\underline{Lal}, \underline{VV}^{1*}$, Mueller, JF^{2} , Aalbersberg, WGL¹

¹Institute of Applied Sciences, the University of the South Pacific, Fiji; ⁴National Research Centre for Environmental Toxicology, the University of Queensland, Brisbane, Australia.

Contact author. Ph: (679) 3232971: <u>lal_v@usp.ac.fj</u>

Introduction

The Stockholm Convention, a global treaty on elimination of persistent organic pollutants (POPs), came into force in May 2004. Fiji was the second country after Canada to ratify the Stockholm Convention, thereby agreeing to formulate a national implementation plan to monitor and control the release of POPs in the country. POPs are hazardous chemicals that are persistent, bioaccumulate in food chains and have the potential to impact at all levels of the trophic system. Fiji is an archipelago of more than 300 islands which lies between $15^{\circ} - 22.5^{\circ}S$ latitude and $174^{\circ}E - 177^{\circ}W$ longitude, spread over a total area of 709,700km² of which 97% is ocean². The current population of Fiji is estimated at about 824,700[°]. The larger islands in Fiji are quite mountainous, but also contain significant flat plains; it is on these flat plains that most agricultural and commercial activities are located and where pollution problems are most likely to occur. Fiji, like many developing countries in the world, has limited or no monitoring data on POPs concentration in the environment and on the exposure of wildlife and humans to these chemicals. Only a few studies on pesticide levels have been carried out in the South Pacific region, including Fiji, and relatively little is known on the occurrence of organochlorine chemicals. The problem is exacerbated by the lack of proper waste management practices, most untreated sewage are discharged into the sea while other waste are typically dumped in open dump sites. Soil and sediment are considered important sinks for dioxin-like chemicals in the environment and their analysis have been used to evaluate potential emission sources of these chemicals. There is very limited scientific data on dioxin (PCDD), furan (PCDF) and dioxin-like PCB (dl-PCBs) concentrations in soil and sediment and sources from the Fiji Islands. In this paper we report concentrations for total dioxin-like compounds (PCDD/F) and

investigate the overwhelming contribution by OCDD congener in all soil and sediment samples taken from various sites representing different land-uses in the Fiji Islands.

Material and methods

Four soil and seven sediment sampling locations representing different land-use types on Fiji Islands were selected to cover a geographical representation of Fiji (Table 1). About 10 cm of the top soil was dug out at each soil sampling site with a clean shovel. Sediment samples were collected from near shore at a water depth of 5 m using a grab sampler made from aluminium. Three replicate subsamples from each site were combined and homogenized. These composite samples were freeze dried, sieved through a 2 mm sieve and placed in individual solvent washed amber jars. Samples were then transported to the National Measurement Institute (NMI) in Sydney, Australia for analysis. Analysis for dioxin-like chemicals (PCDD/PCDF/dl-PCBs) was carried out at NMI, an accredited laboratory using methods adopted from the USEPA Method 1613B and 1668A (ie

isotope dilution technique and quantification using HRMS). Analytes of interest targeted in this study included the 2,3,7,8-chlorine substituted polychlorinated di-benzodioxin and polychlorinated di-benzofuran (PCDD/PCDFs) as well as the $C1_4$ to $C1_7$ PCDD/PCDF homologue groups and dioxin-like PCBs. Recoveries of

the internal/surrogate standard were calculated for all samples. A soil sample used during the 8th round of international intercalibration study (Dioxins 2003) was used as a QCQA sample to assess the methodology

Results and discussions

The soil reference sample, isotopically labelled standards for spiking to calculate recovery and instrument calibration standards were used for quality control and quality assurance purpose. The recoveries of the isotopically labelled internal and surrogate standards were between 70 - 90%. The WHO-TEQ_{DFP} in the Dioxin 6 for the 8th round of international intercalibration study for the sample B soil was 81 pg s⁻¹ dwt (%

2003 report⁶ for the 8th round of international intercalibration study for the sample B soil was 81 pg g⁻¹ dwt (% RSD = 53), the value for WHO₀₅ - TEQ_{DFP} obtained in this study was 62 pg g⁻¹ dry weight (dwt) and was within

that requirement. A number of contaminants including 2,3,7,8-chlorine substituted PCDD/PCDFs as well as the C1_to C1_PCDD/PCDF homologue groups and dioxin-like PCBs were detected.

The concentrations of $\sum PCDD/Fs$ (total of 2,3,7,8-chlorine substituted PCDD and PCDF, C1₄ to C1₇ PCDD/PCDF homologue groups and the dioxin-like PCBs) found in the soil samples from the four sampling locations are summarized in Table 2. The concentration range for $\sum PCDD/F$ detected in the soil samples taken from the various sampling sites was 63 to 5700 pg g⁻¹ (SD = 2700; median = 540) dwt. The current study on soil from different land-use areas in Fiji Islands indicates a dioxin profile where the OCDD is dominant with the corresponding furan detected in lower concentration. The OCDD concentrations contributed 83 - 96% towards $\sum PCDD/Fs$ present in the soil samples from different sampling sites. The highest concentration of OCDD was detected at PU_S02, followed by IN_SO15 (Table 2). At PU_S02, there is a history of chemical misuse (2 tonnes of pesticides were buried and Dicidex was leaking) and at IN_S015, transformer oil was found to have leaked from old transformers⁴. The $\sum PCDD/\sum PCDF$ concentrations ratio (D/F) in soil ranges 13 to 8758. This indicates that OCDD contamination in soil from various land-use types in Fiji is not due to the use of pentachlorophenol (PCP), the D/F ratio for contamination by PCP ranges from 1 – 10⁹. A survey on stockpiles of POPs in Pacific Region highlighted that PCP has been imported for use in Fiji⁴.

The concentrations of \sum PCDD/Fs found in sediment samples from the seven sampling locations are summarized

in Table 2. The concentration range for Σ PCDD/F was 16 to 1000 pg g⁻¹ (SD = 370; median = 63) dwt. The dioxin profile in sediment from all the seven sampling sites shows OCDD concentrations being dominant and contributing 58 - 95% towards Σ PCDD/Fs from different sampling sites. The sediment sample with the highest level of Σ PCDD/F (1000 pg g⁻¹) dwt was from AG_RS20, a site with intensive agriculture especially rice farming and a large scale timber industry. The site (AG_RS20) is also close to a government agricultural research station which acts as a distribution point for pesticides for farmers⁴. The Σ PCDD/ Σ PCDF concentrations ratio (D/F) in sediments ranges 4 to 404. The sites with Σ PCDD/ Σ PCDF concentrations ratio (D/F) within the PCP range included AG_ES11 (4), AG_RS03 (7.3) and PU_MS18 (7.9). However, due to very little contribution by PCDF towards total concentration of PCDD/F it indicates some other source for OCDD abnormality than contamination by PCP¹⁰. Further, 1,2,3,6,7,8-HxCDD is usually present as the dominant isomer among the toxic HxCDDs in PCP¹⁰. The congener 1,2,3,4,7,8-HxCDD is dominant among the 2,3,7,8-substituted HxCDDs in sediments from Fiji Islands.

The similarities in PCDD/F profiles in all soil and sediment from Fiji Islands suggests that PCDD source or formation process in similar in all samples. Further, the OCDD congener represents 73 - 92% towards \sum PCDD/F in various land-use areas in Fiji Islands (Figure 1). This is an interesting find since sampling area covers an area of about 709,700km of which 97% is ocean and includes a variety of anthropogenic and terrestrial influences^{2,11}. Studies in Queensland¹¹, Missisippi in USA¹², Yellow Sea¹³, East China Sea¹³, the Pacific Ocean¹³ and kaolinite samples analysed in Germany¹⁴ have highlighted the dominance of OCDD towards \sum PCDD/F and having similar HxCDD isomer patterns and high \sum PCDD/ \sum PCDF concentration ratios (D/F) in sediment samples. Moreover, these studies highlight that known anthropogenic sources for high concentration of PCDDs (particularly OCDD) were not evident. A widespread source that is natural in origin may be responsible for the OCDD abnormality in soil and sediment samples from Fiji Islands. A number of studies indicate natural formation which include biogenic or geogenic formation processes^{15,16}. Further, environmental processes such as the local climate conditions may be responsible. A faster degradation of the lower chlorinated PCDD/PCDF in the tropical environment may contribute to the observed shift in congener profile. Moreover, the least volatile and most persistent PCDD may accumulate specifically in tropical environments to colder climates where they accumulate⁵. The OCDD dominated dioxin congener profile could also be related to contamination or precursors in pesticides¹⁰ that have been used in Fiji.

Generally, the soil and sediment concentrations of \sum PCDD/F (including OCDD) at all sampling sites were low. Sources of the elevated levels (OCDD) at the peri-urban (PU_S02) and the industrial site (IN_S015) are not known, but the differences between sites can probably be due to local sources. However, the overwhelming representation towards total PCDD/F concentration in all soil and samples across Fiji Islands indicates a widespread source, possibly natural in origin. At this stage, the source or formation process responsible for high concentration of OCDD towards \sum PCDD/F in soils and sediments from various land-use areas in Fiji remains unknown.

Acknowledgments

The authors gratefully acknowledge the provision of a research fellowship to Mr. Vincent Lal for this study by the government of Australia. UNEP/Chemicals for providing technical support in terms of training POPs staff at the Institute of Applied Sciences laboratory in the Fiji Islands during a capacity building project. EnTox is cofunded by Queensland Health.

References

- 1. Aalbersberg WGL., Thaman B. (2000). Institute of Applied Sciences, the University of the South Pacific, Fiji, Suva
- 2. Morrison RJ, Harrison N, Gangaiya P. (1996); Environ Pollut. (93): 159-167.
- 3. Muller J, Muller R, Goudcamp K, Shaw M, Mortimer M, Haynes D, Burniston D, Symons R, Moore M. (2004); *Organohalogen Compd*. (66): 93-99
- 4. Burns T, Graham B, Munro A, Wallis I. (2000); South Pacific Regional Environmental Programme
- Muller J, Muller R, Goudcamp K, Shaw M, Mortimer M, Haynes D, Burniston D, Symons R, Moore M. (2004); National Dioxins Program Technical Report No. 5, Australian Government Department of the Environment and Heritage, Canberra
- 6. Bert van Bavel. (2003). Final Report Eight Round of the International Intercalibration Study
- 7. Holt E, Recke von der R, Vetter W, Hawker D, Alberts V, Kuch B, Weber R. (2008); *Environ Sci. Technol.* (42): 472-1478
- 8.United Nations Environment Programme (UNEP) Chemicals. (2002); Pacific Islands Regional Report. Regional Based Assessment of Persistent Toxic Substances
- 9. Oberg LG, Annerson R, Rappe C. (1992); Organohalogen Compd. (9): 351-354.
- 10.Wittsiepe J, Kullmann Y, Shrey P, Selenka F, Wilhelm M. (1999); *Toxicology Letter Shannon* (106):2-3:191-200
- 11. Gaus C, Papke O, Dennison N, Haynes D, Shaw GR, Conell DW, Muller, JF. (2001); *Chemosphere*. (2001): (43): 549-558
- 12. Hashimoto S, Wakimoto T, Tatsukawa R. (1995); Marine Pollut Bull. (30): 5: 341-346
- Tysklind M, Fangmark I, Marklund S, Lindskog A, Thaning L, Rappe C. (1993); *Environ Sci Technol*. (27):10: 2191-2197
- 14. Rappe C, Anderson R, Bonner M, Kooper K, Fiedler H, Howell F, Kulp SE, Lau C. (1997); *Chemosphere*. (34):1297-1314
- 15. Hagenmaier H, Brunner H. (1987); Chemosphere. (16): 1759-1764
- 16. Jobst H, Aldag R. (2000); Umweltchem. (12):1: 2-4

Sample	Land-use	Sample type	Description
PU_S02	Peri-urban ¹	Soil	Lakena agricultural station
IN_S015	Industrial ²	Soil	Lautoka industrial area
RE_S017	Remote ³	Soil	Monasavu highland (dam)
RE_S019	Remote	Soil	Savusavu (mountain)
AG_RS20	Agricultural ⁴	River sediment	Dreketi agricultural station
AG_ES11	Agricultural	Estuarine sediment	Sigatoka river estuary
RE_MS22	Agricultural	Marine sediment	Levuka harbour
IN_MS05	Remote	Marine sediment	Vatuwaqa induatrial area
AG_RS03	Industrial	River sediment	Wainibokasi jetty, Nausori
PU_MS18	Agricultural	Marine sediment	Savusavu wharf near town
IN_MS06	Industrial	Marine sediment	Suva harbour
IN_SRB	Industrial	Soil	Dioxin200 Intercalibration ⁹

Table 1: Sample nomenclature and description

An area on the fringes of the urban and agricultural boundaries, close to coastal region

An area dominated by industries and is > 5 km from any urban residential population

An area > 60 km from any urban, industrial, agricultural or coastal boundaries

An area > 15 km from any urban settlement, town or cities but close to coastal region

Table 2: Summary of \sum PCDD/F, OCDD and OCDF concentration (pg g ⁻¹ dwt) in soils and sediments					
from various land-use sites studied across Fiji Islands					
	_				

Component (pg g ⁻¹ dwt)	∑PCDD/F	OCDD	OCDF	
Site				
PU_S02	5700	5000	200	
IN_S015	950	910	14	
RE_S017	63	52	3	
RE_S019	120	110	<0.1	
AG_RS20	1000	905	2	
AG_ES11	16	12	< 0.2	
RE_MS22	39	34	1	
IN_MS05	62	59	1	
AG_RS03	19	16	1	
PU_MS18	120	69	10	
IN_MS06	76	65	2	



Figure 1: Percentage (%) contribution by OCDD and OCDF towards ∑PCDD/F in land-use types in Fiji