

EXTRACTABLE ORGANOHALOGENS (EOX) IN BROWN BOOBY (*Sula leucogaster*) FROM NAKANOKAMISHIMA ISLAND, SOUTH RYUKYUS, JAPAN

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Introduction

We have described over five million chemical compounds that exist in our daily life and the chemical industry produces about one hundred and fifty million tons of synthetic chemicals annually ¹. Especially large amounts of organohalogen compounds are produced by the industrial activities of man. Anthropogenic organohalogens are responsible for ecological damage associated with their bioaccumulation and biomagnification potentials and their toxic properties. It is well known that covalently bound halogen atoms, such as chlorine, increase persistency and lipophilicity of an organic molecule, and thus enhance bioaccumulation and biomagnification to toxic levels ¹. However, only a limited number of organohalogen compounds which have been produced in industries have been measured and their occurrence and behaviour determined in the environment ¹. It is not easy to investigate all of the individual compounds and their degradation products due to the cost and time required for analysis. On the other hand, it may be true that there are methodological problems with current measurement techniques for complicated residues of man-made compounds.

Instrumental neutron activation analysis (INAA) is a suitable technique to determine the extent of organohalogen residues in the environment. It is possible to get the levels of individual organohalogen compounds as extractable organohalogen (EOX) concentrations. Analysis of EOX will be of help to understand the levels of unidentified organic halogens when EOX and identified man-made organic halogen concentrations are compared ².

It is well known that determination of the age of wild birds is difficult. Consequently, identification of the age trend of contaminants in wild birds is limited. In the present study, age trend accumulation was investigated for EOX in the seabird samples where age was determined by banding technique.

Materials and Methods

The bird samples were collected at Nakanokamishima Island, South Ryukyus, Japan, during the summer of 1993 with permission from the Japanese government. The five individuals were analysed for EOX and organochlorine compounds [PCBs, DDTs, HCHs, HCB and Chlordane compounds(CHLs)]. Approximately 5 to 10 grams of fat tissue from each sample was used for chemical analysis. EOX and organochlorines were extracted using organic solvents. The extract was washed with distilled deionized water, and fractionated with S-X3 (Bio-Lad Laboratories Inc.) gel permeation column chromatography. EOX in the sample solution was measured by instrumental neutron activation technique^{2,3}. The analysis was made with the research nuclear reactor JRR-4, Japan Atomic Energy Research Institute (JAERI), Ibaragi, Japan, by using neutron flux at a rate of 3.7×10^{13} n/cm² sec for two minutes. The gamma-rays from ³⁸Cl, ⁸⁰Br and ¹²⁸I were measured by gamma-ray spectrometry technique^{2,3}. The reproducibility of this method is 11% for chlorine, 5.0% for bromine and 13% for iodine (n=3). Individual organochlorine compounds such as PCBs, DDTs, HCHs, HCB and CHLs were analysed by GC-MS and GC-ECD⁴. The analytical error of this method was less than 5%.

Results and Discussion

The results of the EOX analysis are shown in Table 1. The EOX concentrations were 56.2 to 111 µg/g (fat weight basis). The order of EOX concentrations was as follows: extractable organochlorine (EOCl) > extractable organobromine (EOBr) > extractable organoiodine (EOI) (Fig. 1). The highest EOCl concentration was found in sample No.120-03889, which was the youngest (three weeks old) and had the lowest lipid content (24.1%) .

The proportion of known organochlorine compounds in EOCl was analysed to reveal the content of unknown organochlorine compounds. The results showed that a large part of EOCl, more than ninety-five percent, was derived from unidentified compounds. This figure is higher than the figure obtained from grey-tailed tattlers(*Heteroscelus brevipes*)⁵. It is observed that brown boobies banded at Nakanokamishima Island travel 1000 to 2000km southward to the southern Philippines⁶. Consequently, this bird species may ingest the compounds via food such as fish and squid⁷ in the regions of southern Japan and the Philippines.

It is already known that some man-made organochlorine compounds, such as PCBs, DDTs, HCHs,

HCB and CHLs, are ubiquitous in the marine environment^{4,8,9}. The results strongly indicated that the contamination of the marine environment includes yet unknown/unidentified organochlorine compounds. The EOCl concentration is especially high in the youngest of the individual sample. It is important to investigate whether these unknown/unidentified compounds cause any physiological effects on brown booby or not, and to determine the sources and the extent of the contamination.

Table 1. Concentrations ($\mu\text{g/g}$ on fat weight basis) of extractable organohalogens (EOX) in five brown boobies (*Sula leucogaster*) from Nakanokamishima Island, South Ryukyus, Japan.

Sample No.	Age (Years)	Fat Content (%)	EOCl	EOBr	EOI
120-03889	0.06	24.1	88	18	5.0
930505-D2	0.06	34.8	40	12	4.6
12A-01132	0.27	58.7	42	12	2.2
120-02366	5.0	55.2	45	11	4.7
120-01785	7.0	52.2	46	12	5.2

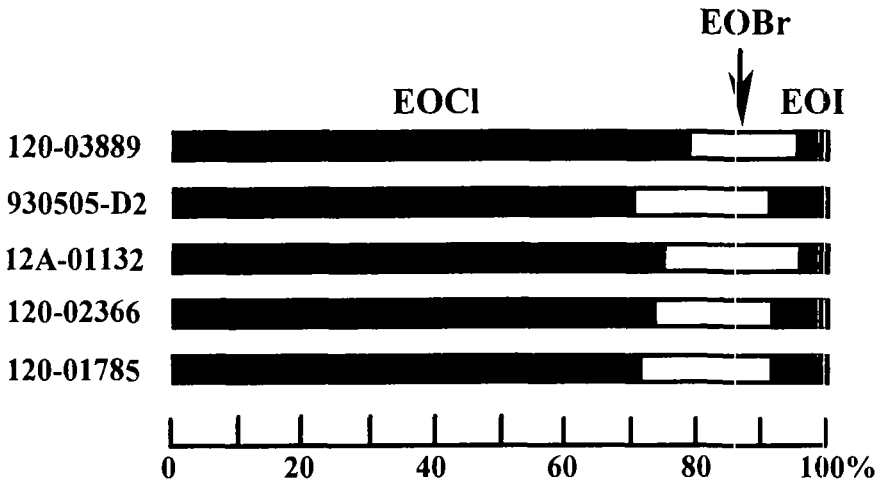


Fig. 1. EOX compositions in adipose tissue of the brown booby samples.

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