

Perfluorinated Sulfonate and Carboxylate Compounds in Seabird Eggs From the Canadian Arctic: Temporal Trends

Birgit M. Braune, Robert J. Letcher

Environment Canada, National Wildlife Research Centre, Carleton University, Ottawa, Ontario, Canada K1A 0H3



Abstract

Temporal trends of perfluorooctane sulfonate (PFOS) and perfluorinated carboxylates (PFCAs) were determined in eggs thick-billed murre and northern fulmars from Prince Leopold Island in the Canadian Arctic from 1975 to 2012. Σ PFCA concentrations increased from 1975 to 2008 in fulmar eggs, and to 2010 in murre eggs, followed by decreases. PFOS concentrations did not change significantly. PFUnA (C₁₁) and PFTrA (C₁₃) were the major PFCAs.

Introduction

- Poly- and per-fluorinated alkyl substances (PFASs) are ubiquitous in the Arctic environment.
- The major fluorinated compounds measured are the perfluorinated sulfonates (PFSAs) - e.g. perfluorooctane sulfonate (PFOS) - and the PFCAs which include perfluorooctanoate (PFOA).
- PFASs can biomagnify and have been found in arctic biota including seabirds and their eggs.
- In this study, we examined temporal trends of PFOS and PFCAs in eggs of two seabird species, the thick-billed murre (*Uria lomvia*) and northern fulmar (*Fulmarus glacialis*), from the Canadian Arctic.

Methods

- From 1975 to 2012, eggs of northern fulmars and thick-billed murre were collected from Prince Leopold Island (74° 02'N, 90° 05'W) in Lancaster Sound.
- Egg homogenates were analyzed as pooled (composite) samples consisting of three eggs.
- The PFAS extraction, cleanup and analysis have been described in Chu and Letcher (2008) and Gebbink et al. (2011).
- PFCAs and PFSAs were determined based on liquid chromatography-negative electrospray ionization-tandem mass spectrometry (LC-ESI(-)-MS/MS).
- Quantification used an internal standard approach; concentrations were inherently recovery-corrected.

Data Treatment

- Since PFASs such as PFCAs and PFSAs bind to proteins rather than partition into lipid, concentrations were not lipid-normalized.
- Statistical tests were performed using Statistica for Windows Version 7.0 with a significance level of $p < 0.05$.
- Temporal trends were analyzed by backward stepwise regression analysis, with year and $\delta^{15}\text{N}$ as regressors.

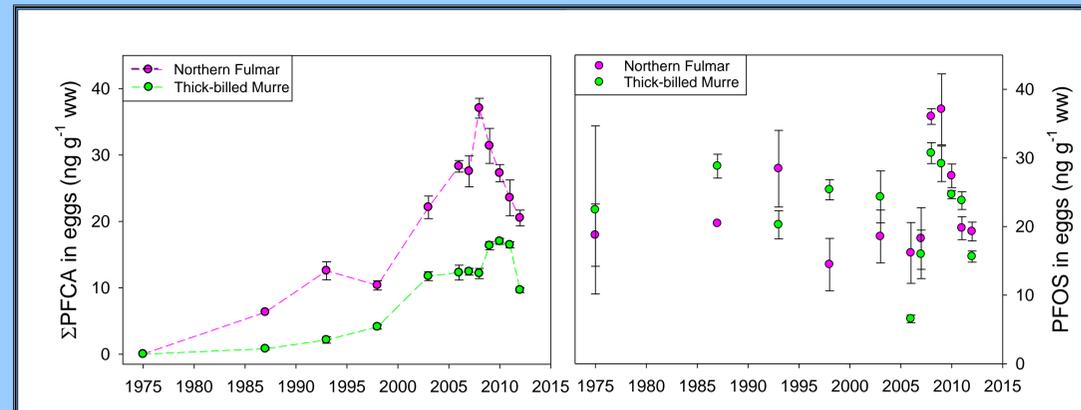


Figure 1. Mean annual concentrations (\pm standard error) of Σ PFCA and PFOS in eggs of northern fulmars and thick-billed murre from Prince Leopold Island, Nunavut, Canada, 1975-2012. Σ PFCA = sum of PFHxA (C₆), PFHpA (C₇), PFOA (C₈), PFNA (C₉), PFDA (C₁₀), PFUnA (C₁₁), PFDaA (C₁₂), PFTrA (C₁₃), PFTeA (C₁₄) and PFPA (C₁₅).

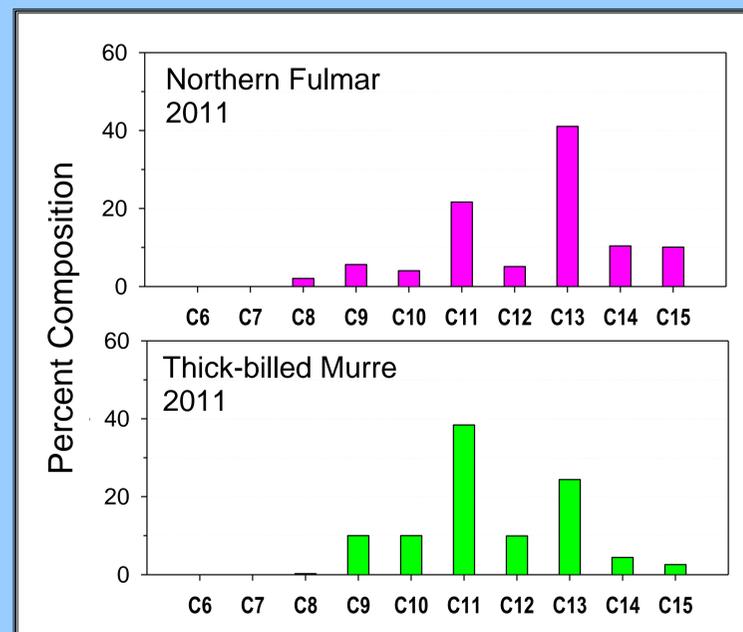


Figure 2. Mean contributions (%) PFHxA (C₆), PFHpA (C₇), PFOA (C₈), PFNA (C₉), PFDA (C₁₀), PFUnA (C₁₁), PFDaA (C₁₂), PFTrA (C₁₃), PFTeA (C₁₄) and PFPA (C₁₅) to Σ PFCA in eggs of northern fulmars and thick-billed murre from Prince Leopold Island, Nunavut, Canada, 2011.

Results

- Σ PFCA levels increased significantly from 1975 to 2010 in northern fulmar eggs ($n=47$, $r=0.89$, $p<0.0001$) and from 1975 to 2008 in murre eggs ($n=35$, $r=0.88$, $p<0.0001$) followed by decreases (Figure 1, left graph).
- PFOS has not changed significantly although levels may be decreasing in recent years (Figure 1, right graph).
- Significant increases in concentrations of all longer-chained PFCAs (C₉-C₁₅) contributed to the increase in Σ PFCA in the murre and the fulmars.
- PFUnA (C₁₁) and PFTrA (C₁₃) were the predominant PFCAs measured in eggs of both species (Figure 2). These two PFCAs together constituted $>60\%$ of Σ PFCA in all years, with PFTrA dominating the fulmar PFCA profile and PFUnA dominating the murre PFCA profile.
- PFOA (C₈) has been detected in eggs of both species only since 2008 and comprises $<3\%$ of the PFCA profile.
- In 2011, the fulmar PFCA profile showed a stronger presence of the longer-chained PFCAs (C₁₀-C₁₅) which comprised $>90\%$ of Σ PFCA, whereas in the murre, the profile was dominated ($>90\%$) by the C₉-C₁₃ PFCAs (Figure 2).

Acknowledgements

- Egg collections: D. Nettleship, A. Gaston, M. Mallory and field crews.
- Sample processing: Laboratory Services personnel, National Wildlife Research Centre (NWRC), Ottawa.
- Chemical analyses: S. Chu and D. Blair, OCRL, NWRC, Ottawa.
- Stable isotope analyses: Keith Hobson, Environment Canada.
- Funding: Environment Canada; Northern Contaminants Program of Aboriginal Affairs and Northern Development Canada.
- Logistical support: Polar Continental Shelf Project, Natural Resources Canada.

Literature Cited

- Chu SG, Letcher RJ. 2008. *J. Chromatogr. A* 1215: 92-99.
 Gebbink WA, Letcher RJ, Hebert CE, Weseloh DVC. 2011. *J. Environ. Monit.* 13: 3365-3372.



Environment Canada
 Environnement Canada