Total mercury in hair of polar bears (*Ursus maritimus*) from Greenland and Svalbard

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Concentrations (ppm = μg/g dry weight) of total mercury (Hg) were determined in hair of polar bears (*Ursus maritimus*) from northwestern Greenland (N = 22; period of sampling: 1978-1989), eastern Greenland (N = 44; 1984-1989) and Svalbard (N = 31; 1980). For subadults (2-6 years of life), adults (7-10 years), and old bears (>10 years), concentrations of total Hg in hair were not found to be dependent on age or sex. A decreasing trend in Hg concentrations was found from west to east. The mean concentrations of total Hg in hair (cubs of the year and yearlings excluded) were: northwestern Greenland, \( x = 8.38 \) ppm (min.-max.: 1.71-14.19 ppm, N = 21); eastern Greenland: \( x = 4.58 \) ppm (min.-max.: 2.50-8.83 ppm, N = 41); and Svalbard, \( x = 1.98 \) ppm (min.-max.: 1.02-4.55 ppm, N = 29). Concentrations found in northwestern Greenland were similar to those reported by others from the hair of polar bears sampled within management zone F of the eastern Canadian High Arctic. Concentrations of total Hg in polar bear hair from eastern Greenland were similar to concentrations found by others in contemporary (1988) material collected during spring in western Svalbard. However, the mean concentration of total Hg in the 1980 Svalbard material, which was collected during July-September, was significantly lower than concentrations found in samples taken during late winter and spring in eastern Greenland and at Svalbard, respectively. Presumably the relatively low concentrations found in the 1980 Svalbard sample are attributable to the period of moult and hence a larger proportion of newly grown hair in the individual samples. In a subsample consisting of internal tissues from 19 polar bears from eastern Greenland (1984-1987), concentrations of total Hg in hair correlated positively with concentrations of total Hg (wet weight) in muscle (N = 6), liver (N = 19) and kidney (N = 19) tissue. For liver and kidney tissue these relationships were statistically significant.

**Introduction**

Hair samples of mammals are considered to be good indicators of the body burden of mercury (Jóhannesson et al. 1981; Hansen 1981; Renzoni 1989). Concentrations in hair reflect blood mercury or body burden of mercury during time of hair growth (Sexton et al. 1978; Kershaw et al. 1980). Being a top predator which feeds almost exclusively on seals and in particular ringed seals (*Phoca hispida*; e.g. Stirling & Archibald 1977; Lønne 1970), the polar bear can be considered a good indicator of the degree of bioaccumulation of persistent contaminants in its ecosystem. Based on analyses of 128 recent (1977-1980) and 18 museum samples (1910-1927), Eaton & Farant (1982) presented concentrations of total mercury (Hg) in polar bear hair from eight different localities in the Canadian High Arctic. Liver samples from 67 polar bears from 1982 were analysed for mercury and 21 other elements by Norstrom et al. (1986). Renzoni & Norstrom (1990) gave concentrations of total Hg in hair sampled between 1976 and 1988 from a total of 141 polar bears in various localities in the Canadian Arctic, the USSR, and at Svalbard (Norway). Concentrations of total and organic Hg in muscle, liver and kidney in polar bears from central-eastern Greenland were presented in Dietz (1987), Dietz et al. (1990) and Dietz & Agger (in press). Here we present the results of analyses of total Hg in hair from 66 polar bears from three localities in Greenland with a note on the relationship between concentrations of total Hg found in hair and muscle, liver and kidney tissue. For comparative reasons analyses of total Hg in 31 polar bear hair samples obtained at Svalbard during July-September 1980 are included.
Material and methods

A total of 66 hair samples (28 F, 37 M, 1 undet.) from Greenland and 31 samples (15 F, 16 M) from Svalbard have been analysed for contents of total Hg. The majority of the Greenland samples were collected by hunters from their subsistence catch of polar bears. Between January 1984 and August 1989 a total of 40 samples (year/sample size: 1984/11; 1985/2; 1986/1; 1987/4; 1988/8; 1989/13; unknown/1) were obtained from the Scoresby Sound region of eastern Greenland (Fig. 1). Four samples (2F, 2M) were obtained in 1989 from the Ammassalik area in southeastern Green-

![Map of polar bear sample locations](image-url)

*Fig. 1. Areas in which polar bear hair samples analysed in this study were obtained. The borders of the Canadian polar bear management zone F are shown with a dotted line (---).*
land; a total of 22 samples (8 F, 14 M) were collected in the Avanersuaq (Thule) area in northwestern Greenland. Four of these samples were collected by scientists in May 1978; the remainder were collected by local hunters in 1988 (N = 7) and 1989 (N = 11). A total of 31 hair samples were collected by scientists during a Swedish icebreaker expedition (YMER) operating in the Svalbard area in 1980. The vast majority of the Greenland samples were from late winter and spring (January–May) while the Svalbard-material was collected during “summer” (July–September; Fig. 2). For 19 of the polar bears from the Scoresby Sound material (1984–1987), additional information was available on total mercury content in muscle (N = 6), liver (N = 19) and kidney (N = 19) tissue.

The hair samples were kept frozen in polyethylene plastic bags until analysis (with the exception of the 1978 Thule samples, which were stored dry in paper bags). The samples were carefully washed in acetone, rinsed and dried before being analysed for mercury residues at the laboratory of Dipartimento Biologia Ambientale (Siena) by atomic absorption spectrophotometry as described in Renzoni et al. (1986). Samples of internal tissues were kept frozen and stored in polyethylene plastic bags before being analysed at the laboratory of Greenland Environmental Research Institute by a method described in Dietz et al. (1990).

To detect whether concentrations of total Hg in hair correlated with age, the polar bears were grouped into five age classes: cub of the year = COY, yearling, subadult, adult, and old. The age classes were based either on age obtained from reading of the annual layering in tooth cementum or on field age class estimates and body length. For a sub-sample from Scoresby Sound (N = 19; 1984–1987), individual ages were determined from the counting of the annual layering in the cementum of the lower premolars as described in Stirling et al. (1977). For the 1978 Thule-sample and the 1980 Svalbard-sample, ages were esti-

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Fig. 2. Seasonal distribution of polar bear sampling from three different regions in Greenland (Thu = Thule; Sco = Scoresby Sound; Amm = Ammassalik) and Svalbard (Sva). Four samples from Scoresby Sound presented in Table 2 had no information on the date of sampling.
mated in the field by scientists. The majority of the Greenland material was categorized into age classes according to estimates of age given by the hunters in the field together with zoological body length measured from tip of snout to tip of tail along the back when the bear was positioned on its belly. Length data for these age classes are presented in Table 1 where adult and old females are pooled because there was no difference in average body length of these age classes (F = 0.595, df = 1/18, P > 0.05).

Concentrations of total Hg are given as ppm (µg/g. d.w.) in hair and as ppm (µg/g. w.w.) in other tissues.

The distributions of Hg in hair in the different areas did not differ significantly from normality (Kolmogorov-Smirnov one-sample tests; P > 0.05) but had significantly different variances (Bartlett’s test, X² = 19.567, P < 0.01). All data were ln-transformed and ANOVA analyses were performed using the factors: area, sex, age, and the respective interactions of these. Statistics were made using Statview II and SuperANOVA for a Macintosh microcomputer.

### Results

#### Hg in hair

Concentrations of total Hg found in polar bear hair in the four different areas are presented in Table 2. Values for the two youngest age groups are given separately because young bears (i.e. cubs of the year = COYS and yearlings) are assumed to be dependent on milk until they are well into their second year of life (Lømo 1970). Most cubs become independent when they are little over 24 months old (Larsen 1986). In none of the samples were concentrations of total Hg in hair related to either age (age classes: subadults, adults, olds) or to sex (two factor ANOVAs; P > 0.05). Therefore, data from these two categories were pooled for further analyses. As concentrations of total Hg in polar bear from Ammassalik in southeastern Greenland did not differ significantly from concentrations found in hair from the Scoresby Sound region (F = 0.446, P < 0.05. df = 1/39), the samples from these

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### Table 1. Polar bear age classes with zoological body lengths used in analyses of relationship between total Hg in hair and age

| Age class       | Mean length (cm) | Min.-Max. | N |
|-----------------|------------------|-----------|---|
| COY M           | 107              | -         | 1 |
| F               | -                | -         |   |
| Yearlings M     | 177              | -         | 1 |
| F               | 143              | 13.2-157  | 4 |
| Subadults M     | 200              | 16.9-173  | 14|
| (2-6 yr) F      | 140              | 18.2-223  | 16|
| Adults (7-10 yr)M | 224             | 19.5-215  | 15|
| Old (10 yr) M   | 242              | 18.1-273  | 17|
| Adults and old (7 yr and older) F | 213 | 16.6-176-220 | 20|

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### Table 2. Concentrations (µg/g. d.w.) of total mercury in hair of polar bears from Greenland (1978-1989) and Svalbard (1980). The average and error ranges are calculated from the antilogarithms of the mean and the standard deviation of the ln-transformed data.

| Location   | Year     | Age groups       | Geo. Mean | Standard error range | Min.-Max. | N  |
|------------|----------|------------------|-----------|----------------------|-----------|----|
|            |          |                  | e^±SD-e^-SD |                      |           |    |
| Thule      | 1978-1989| Yearling         | 9.51      | -                    | -         | 1  |
|            |          | 2 years and older| 8.38      | 6.38-11.01           | 4.71-14.19| 21 |
| Scoresby Sound | 1984-1989| Yearlings       | 1.81      | 1.63-2.01            | 1.65-2.03 | 3  |
|            |          | 2 years and older| 4.62      | 3.39-6.17            | 2.50-8.83 | 37 |
| Ammassalik | 1989     | 2 years and older| 4.21      | 3.68-4.82            | 3.41-6.02 | 4  |
| Svalbard   | 1980     | Cub of the year  | 0.34      | -                    | -         | 1  |
|            |          | Yearling         | 1.04      | -                    | -         | 1  |
|            |          | 2 years and older| 1.98      | 1.33-2.96            | 1.02-4.55 | 20 |
areas were pooled. The geometric mean concentrations of Hg differed significantly between the three main areas (F = 122.09, df = 2/88, P < 0.01). The northwestern Greenland Hg-concentration was 1.8 times and 4.3 times higher than mean concentration in eastern Greenland and at Svalbard, respectively (Table 2).

Concentrations found in hair from eastern Greenland were significantly higher than concentrations found in the sample taken at Svalbard in 1980, but similar to concentrations reported by Renzoni & Norstrom (1990) for a sample obtained from Svalbard during the spring of 1987 and 1988 (Tukey's test; P < 0.01).

In the East Greenland sample the concentrations of total Hg in hair collected during July and August (X = 5.23 ppm, rel. SD = 1.26, min.-max.: 3.54–6.62 ppm, 5M and 1F) did not differ significantly from concentrations in hair at other times of the year (F = 0.066; df = 3/33; P < 0.05). Insufficient seasonal representation did not allow similar analyses to be performed for the other samples.

Hg in muscle, kidney and liver

The relationships of concentrations of total Hg in hair to concentrations of total Hg in muscle, kidney and liver tissue from 19 polar bears sampled in the Scoresby Sound region (eastern Greenland) between 1984 and 1987 in the period December–April are shown in Fig. 3. Concentrations of mercury in hair were found to be positively correlated with concentrations in tissue from kidney (r = 0.480, N = 19, P < 0.05), liver (r = 0.685, N = 19, P < 0.05), and muscle (r = 0.574, N = 6, n.s.). The slopes of the regression lines were significantly different from 0 at the same probability level for kidney and liver tissue, whereas no significance was detected for muscle tissue, probably due to the low number of samples.

Discussion

Concentrations of total Hg in polar bear hair were highest in northwestern Greenland. However, the concentrations were in the same order of magnitude as those reported by Eaton & Farant (1982) and Renzoni & Norstrom (1990) from the eastern Canadian High Arctic (i.e. within the Canadian management zone F; Fig. 1). Recaptures in northwestern Greenland Hg-concentration was 1.8 times and 4.3 times higher than mean concentration in eastern Greenland and at Svalbard, respectively (Table 2).
western Greenland of polar bears which were marked in zone F (Rosing-Asvid & Born 1990) indicate that polar bears move freely between these areas.

Concentrations of total Hg in hair from polar bears from eastern Greenland were similar to values reported by Renzoni & Norstrom (1990) for a sample taken during spring in western Svalbard. Satellite tracking (Larsen et al. 1983) and movements of marked bears (Larsen 1986) indicate that exchange occurs between polar bears in eastern Greenland and in the Svalbard – Frans Josef Land region. However, to what extent local groups of bears exist within this range still remains undetermined (Born & Rosing-Asvid 1989).

Eaton & Farant (1982) and Renzoni & Norstrom (1990) suggested that geographical variation in concentrations of total Hg in the hair of polar bears in the Canadian High Arctic to some extent reflects different ratios of the bottom-feeding bearded seals (Erignathus barbatus) to the more pelagic ringed seals in the diet of the bears in the various areas. The very limited and highly heterogeneous information available on the feeding of polar bears in Greenland (Pedersen 1945; Born upubl. data) and at Svalbard (Lone 1970) does not allow for an evaluation of geographical variation in polar bear diet in these areas. On the other hand, Dietz & Agger (in press) state that the only differences found in mercury levels in bearded and ringed seals from NW Greenland are found in liver tissue, as also found in Smith & Armstrong (1975) in Canada, whereas no differences were found between these two species in blubber and muscle and kidney tissue.

We suggest that the differences found in our study in concentrations of total Hg in hair from Greenland polar bears mainly reflect variations in the overall mercury burden of the environment rather than different proportions of various prey species in their diet.

Norstrom et al. (1986) found that mercury concentrations in polar bears from the western part of the Canadian Arctic tended to be higher than in the eastern part. Likewise mercury concentrations in the northern areas at Melville Island were higher than in southern regions. Dietz & Agger (in press), who found that concentrations of total Hg in the liver tissue of polar bears from eastern Greenland were lower than those reported by Norstrom et al. (1986), confirm a decreasing trend in Hg concentrations from west to east.

Higher concentrations of Hg were found in polar bear hair sampled at southwestern Svalbard in 1987 and 1988 (Renzoni & Norstrom 1990) than in a sample from the pack ice areas around the archipelago in 1980. To our knowledge there is no information which indicates that the total mercury burden of the marine environment of the Svalbard region has increased within the last decade and we suspect that the relatively low concentrations of total Hg found in the 1980-sample from Svalbard reflect seasonal variation related to moulting of hair. The 1980 Svalbard-samples (this study) were collected from “summer fur” while the samples analysed by Renzoni & Norstrom (1990) were taken during March–April from “winter fur.” The majority of samples from Greenland were collected between January and May from “winter fur” (Fig. 2) which presumably is also the case with samples in other studies (Eaton & Farant 1982; Renzoni & Norstrom 1990). Although information on the timing of the process of moulting in polar bears is somewhat contradictory (Freuchen 1935; Pedersen 1945; Uspenski 1939; Vibe 1981), it appears that generally the moult begins in late May and that the new winter coat has grown in autumn (September–October?).

Our assumption that hairs of “summer fur” have lower concentrations of total Hg than hairs of winter fur was, however, not supported by the finding that in only six samples, taken in July and August in eastern Greenland, the mean concentration was not significantly different from the mean concentration in hairs sampled in the same area during other seasons.

We suggest that the low Hg levels found in “summer” hair are caused by accelerated hair growth during the moult. However, it remains undetermined whether the different concentrations found at Svalbard (this study and Renzoni & Norstrom 1990) also reflect sampling of different segments of the polar bear population at different times of the year and/or seasonal variations in the food of polar bears.

In samples from 55 polar bears taken between January and May in 1982 in seven areas within the central Canadian High Arctic, Renzoni & Norstrom (1990) did not find a statistically significant correlation between total Hg in hair and live tissue. In contrast, in a smaller sample taken during spring in eastern Greenland we found that concentrations of total Hg in hair correlated positively with concentrations of total Hg in internal
tissues with a capacity for long term accumulation of mercury. An explanation for this apparent discrepancy found in the two studies could be that samples from various areas in Canada were pooled in the Renzoni & Norstrom (1990) study whereas the East Greenland material is more homogeneous since it was sampled in only one area.

The mercury concentrations in the hair as well as in the internal organs reflect the contamination levels in the food of the polar bears. Therefore, it appears less surprising that we found a correlation between Hg in hair and in internal tissues. This is in agreement with previous findings by for example Freeman & Horne (1973), who stated that claws and fur from the Gray seal (Halichoerus grypus) and the harp seal (Pagophilus groenlandicus) were good indicators of the degree of mercury contamination in muscle, liver, kidney and heart tissue. Similarly, Bacher (1985) found a significant correlation between mercury in hair and in internal tissues (muscle, liver, spleen, brain) of Australian fur seals (Arctocephalus pusillus). Furthermore, if one regards hair as one of the excretion routes of Hg (see Itano et al. 1984), it seems reasonable to assume that Hg concentrations in hair in a general way reflect the Hg level in the internal tissues of an individual.

We conclude that concentrations of total mercury in polar bear hair from late winter and spring reflect the overall mercury burden in the ecosystem within the range of the various polar bear populations. We suggest that the relatively low concentrations of total mercury found in samples obtained during summer can be related to the process of moulting. Based on a small sample from eastern Greenland we have found indications that concentrations of total Hg in hair taken during spring reflect levels of total mercury in liver and kidney tissue.

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