Are liver and renal lesions in East Greenland polar bears (Ursus maritimus) associated with high mercury levels?

Sonne, Christian; Dietz, Rune; Leifsson, Páll Skúli; Asmund, Gert; Born, Erik W.; Kirkegaard, Maja

Published in:
Environmental health

DOI:
10.1186/1476-069x-6-11

Publication date:
2007

Document version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
Sonne, C., Dietz, R., Leifsson, P. S., Asmund, G., Born, E. W., & Kirkegaard, M. (2007). Are liver and renal lesions in East Greenland polar bears (Ursus maritimus) associated with high mercury levels? Environmental health, 6(11). https://doi.org/10.1186/1476-069x-6-11
Are liver and renal lesions in East Greenland polar bears (*Ursus maritimus*) associated with high mercury levels?

Christian Sonne*1, Rune Dietz1, Pall S Leifsson2, Gert Asmund1, Erik W Born3 and Maja Kirkegaard1

Address: 1Section for Contaminants, Effects and Marine Mammals, Department of Arctic Environment, National Environmental Research Institute, University of Aarhus, Frederiksbergvej 399, POBox 358, DK-4000 Roskilde, Denmark, 2Department of Veterinary Pathobiology, Faculty of Life Sciences, University of Copenhagen, Bülowsvej 17, DK-1870 Frederiksberg, Denmark and 3Greenland Institute of Natural Resources, P.O. Box 570, DK-3900 Nuuk, Greenland

Email: Christian Sonne* - csh@dmu.dk; Rune Dietz - rdi@dmu.dk; Pall S Leifsson - ple@life.ku.dk; Gert Asmund - gas@dmu.dk; Erik W Born - ewb@dpc.dk; Maja Kirkegaard - mki@dmu.dk

* Corresponding author

Abstract

**Background:** In the Arctic, polar bears (*Ursus maritimus*) bio-accumulate mercury as they prey on polluted ringed seals (*Phoca hispida*) and bearded seals (*Erignathus barbatus*). Studies have shown that polar bears from East Greenland are among the most mercury polluted species in the Arctic. It is unknown whether these levels are toxic to liver and kidney tissue.

**Methods:** We investigated the histopathological impact from anthropogenic long-range transported mercury on East Greenland polar bear liver (n = 59) and kidney (n = 57) tissues.

**Results:** Liver mercury levels ranged from 1.1–35.6 µg/g wet weight and renal levels ranged from 1–50 µg/g wet weight, of which 2 liver values and 9 kidney values were above known toxic threshold level of 30 µg/g wet weight in terrestrial mammals. Evaluated from age-correcting ANCOVA analyses, liver mercury levels were significantly higher in individuals with visible Ito cells (p < 0.02) and a similar trend was found for lipid granulomas (p = 0.07). Liver mercury levels were significantly lower in individuals with portal bile duct proliferation/fibrosis (p = 0.007) and a similar trend was found for proximal convoluted tubular hyalinisation in renal tissue (p = 0.07).

Background

Approximately 200–300 tons mercury (Hg) is yearly transported to the Arctic from lower latitudes via the atmosphere and large scale ocean currents [1,2], hence the Arctic acts as a sink for global emitted mercury due to spring mercury depletion events [3-6]. Mercury is toxic and the organic methyl-mercury (MeHg) form mediates central nervous system toxicity pre- and postnatally mainly via the posterior cortex [7,8]. Mercury levels are high in the Greenland and Faroe populations and a special focus has been to address health effects in these regions as such effects are proposed to have a significant socioeconomic impact [9].

The inorganic form of mercury is toxic to liver and kidney tissues due to the co-enzyme inhibition via high affinity to various microsome and mitochondria SH-group enzymes [8]. Studies of Atlantic bottlenose dolphins (*Tursiops truncatus*) [10] and Arctic beluga whales (*Delphinapterus leucas*) [11] have linked mercury exposure to liver...
histopathological changes. Also, renal glomerular and tubular lesions have been associated with mercury toxicity in controlled laboratory studies [12-14], in humans [15,16], in wildlife [11,17] and in domestic mammals [18,19].

In the Arctic, polar bears (Ursus maritimus) bio-accumulate mercury as they prey on polluted ringed seals (Phoca hispida) and bearded seals (Ergignathus barbatus) which results in a high mercury uptake [1,2,20]. Studies have shown that East Greenland polar bears are among the most mercury polluted species in the Arctic [1,2]. In target organs like liver and kidney, inorganic mercury concentrations of 2.13–13.4 and 2.87–32.0 µg/g w.w. (wet weight) have been reported [21,22]. These levels exceed threshold levels for lethal mercury toxicity [1,2]. Although mercury is a naturally occurring element in the Arctic, Dietz et al. [23] showed that up to 94% of the mercury in East Greenland polar bears is of anthropogenic origin. We therefore examined liver and kidney histology and mercury concentrations in East Greenland polar bears sampled during 1999 to 2002 to determine whether high mercury concentrations may cause pathological changes at the histological level in target organs of large Arctic top predators.

Methods

Sampling
Liver (n = 59) and kidney (n = 57) samples were taken by local subsistence hunters in the Scoresby Sound area in central East Greenland (69°00’N to 74°00’N) during 1999–2002. A randomly chosen single renal lobe and a tissue sample from the periphery of a right liver lobe was taken for histological examination and fixed in a phosphate buffered formaldehyde/alcohol solution (3.5% formaldehyde, 86% ETOH and 10.5% H2O) to avoid freeze damage. In addition, sub samples for mercury analyses were stored in separate Polyethylene plastic bags until arrival at the laboratory in Roskilde. All samples were taken <12 h post mortem.

The sample
The sample consisted of 32 subadults (10 females and 22 males), 15 adult females (5 ≥ 15 years) and 12 adult males (4 ≥ 15 years). Of these, 32 were caught during summer (1 June to 30 September) and 27 during winter (1 October to 31 May). The general East Greenland polar bear liver and renal histology, including micrographs, is described by Sonne et al. [24,25].

Age estimation
The age determination was carried out by counting the cementum Growth Layer Groups (GLGs) of the lower I3 tooth after decalcification, thin sectioning (14 µm) and staining (toluidine blue) using the method described by Dietz et al. [26]. Adult males were categorised as ≥ 6 years, adult females as ≥ 5 years, old as ≥ 15 years and the remaining as subadults [27,28]. Furthermore, seasonal differences were investigated as summer: 1 June to 30 September vs. winter: 1 October to 31 May [27].

Liver histology
The liver tissue was trimmed, processed conventionally, embedded in paraffin, sectioned at about 4 µm and stained with Haematoxylin (Al-Haematein)-Eosin (HE) and periodic acid-Schiff (PAS) for routine diagnostics, Van Gieson and Masson Trichrome to detect fibrous tissue (collagen), Best’s carmine to demonstrate glycogen storage, Sudan III to detect lipid (frozen tissue) and Perls’ Prussian blue reaction and Smorl for detecting hemosiderin and lipofuscin pigments, respectively [29,30].

Six histological features were evaluated: 1. Portal mononuclear cell infiltrations (absent, unifocally, multifocally or diffuse). 2. Random mononuclear cell infiltrations (average no. in 5 fields at 10× magnification). 3. Lipid granulomas (average no. in 5 fields at 10× magnification). 4. Hepatocytic steatosis (intracellular lipid; absent, foamy, multifocally macrovesiculary or diffuse macrovesiculary). 5. Visible Ito cells (average no. in 5 fields at 20× magnification). 6. Mild multifocally bile duct proliferation accompanied by portal fibrosis (absent or present). Each histological change was grouped semi-quantitatively as: 1. Portal mononuclear cell infiltrations: absent = 0; mild = unifocally; moderate = multifocally and severe = diffuse. 2. Random cell infiltrations: absent = 0; mild = Index [0;1]; moderate = Index [1;3] and severe = Index [3;8]. 3. Lipid granulomas: absent = 0; mild = Index [0;1]; moderate = Index [1;2] and severe = Index [2;5]. 4. Hepatocytic steatosis: mild = foamy; moderate = multifocally macrovesiculary and severe = diffuse macrovesiculary. 5. Ito cells: absent = 0; mild = Index [0;10]; moderate = Index [10;50] and severe = Index [50;200]. 6. Bile duct proliferation; see above.

Renal histology
Kidney tissue was trimmed, processed conventionally, embedded in paraffin, sectioned at about 4 µm and Periodic acid-Schiff (PAS) and periodic acid silver methenamine (PAS-M) were used to demonstrate glomerular (capillary and mesangial) and proximal convoluted tubular changes; Van Gieson and Masson Trichrome to detect fibrous tissue (collagen) in the glomeruli (glomerulofibrosis) and in the interstitium (interstitial fibrosis) [30]. Seven histological features were evaluated: 1. Glomerular capillary wall thickening (10 randomly selected 5–40× fields). 2. Glomerular mesangial deposits (10 randomly selected 5–40× fields). 3. Tubular epithelial cell hyperplasia (10 randomly selected 5–40× fields). 4. Proximal convoluted tubular hyalinization/atrophy/dilations/necrosis (10 randomly selected 5–40× fields). 5. Tubular medullar

Environmental Health 2007, 6:11
http://www.ehjournal.net/content/6/1/11
hyaline casts (10 randomly selected 5–40× fields). 6. Interstitial fibrosis (10 randomly selected 5–40× fields). 7. Mononuclear cell infiltrations (10 randomly selected 5–40× fields). Each histological change was grouped semi-quantitatively as: 1. Glomerular capillary wall thickening: absent, mild and moderate. 2. Glomerular mesangial deposits: absent, mild and moderate. 3. Tubular epithelial cell hyperplasia: mild: focally; moderate: multi focally. 4. Proximal convoluted tubular hyalinization/atrophy/dilatations/necrosis: mild: focally; moderate: multi focally. 5. Tubular medullar hyaline casts: mild: focally; moderate: multi focally. 6. Interstitial fibrosis: mild: focally; moderate: multi focally. 7. Mononuclear cell infiltrations: mild: focally; moderate: multi focally.

Analyses of mercury

Liver (n = 59) and kidney (n = 57) samples were analysed for mercury levels (µg/g w.w.) according to Dietz et al. [23]. The principle was atomic absorption spectrometry (AAS; hydride generation and the flow injection analyses) have previously been described by Asmund et al. [31]. The detection limit was 0.005 mg/kg of dry weight. Analytical quality was ensured by repeated analyses and by frequent analysis of various certified reference materials [TORT-2 (lobster hepatopancreas), DORM-2, and Dolt-3] supplied by the National Research Council of Canada (Marine Analytical Chemistry Standards Program).

Statistical analyses

The statistical analyses were performed with the SAS statistical software package (SAS V8 and enterprise guide V1) and the level of significance was set to p ≤ 0.05, while significance at 0.05 < p ≤ 0.10 was considered a trend. The mercury data were log-transformed (base e) prior to the analyses in order to meet the assumption of normality and homogeneity of the variance.

For each specific histologic change, a One-way ANOVA was performed to test for differences in mean age between individuals with and without that specific histologic change (Table 1). In case of hepatocytic steatosis, foamy cytoplasm was tested against macro vesicular lipid accumulation. Furthermore, we tested whether there was a relationship between sex and season ([summer: 1 June to 30 September]; [winter: 1 October to 31 May]), respectively, and histologic lesions using a χ²-test. In case of age dependency, a χ²-test was performed within subadults, adults and old, respectively, to test for sex differences.

The difference between mercury concentration and sex was tested in an age-normalizing analysis of covariance (ANCOVA) with mercury concentration as the dependent variable, age as covariable and sex as class variable, including their 1st order interaction links (age × sex). After a successive reduction of non-significant interactions, judged from the type-III sum of squares (p ≤ 0.05), the significance of each of the remaining factors was evaluated from the final model Least Square Mean (LSMean). Then, a One-way ANOVA was performed to test for differences in mercury mean concentration between subadults, adult females and adult males. The results were finally evaluated from Tukey’s post hoc test (Table 2). In order to test the relationship between concentrations of mercury and age, a linear regression model was employed for subadults, adult females and adult males, respectively (Table 2).

### Table 1: Prevalence [% [n]] of liver and kidney histopathology in East Greenland polar bears sampled during 1999–2002

| Organ | Absent | Mild | Moderate | Severe | Age p (F) |
|-------|--------|------|----------|--------|-----------|
| Liver (n = 59) | | | | | |
| Multifocally mononuclear cell infiltrations | 89 (53) | 11 (6) | - | - | n.s. |
| Portal mononuclear cell infiltrations | 85 (50) | 8 (5) | 7 (4) | - | n.s. |
| Lipid granulomas | 37 (22) | 36 (21) | 22 (13) | 5 (3) | n.s. |
| Hepatocytic steatosis | - | 24 (14) | 24 (14) | 52 (31) | n.s. |
| Ito cell lipid accumulation | 36 (21) | 14 (8) | 19 (11) | 31 (19) | *** (7.8) |
| Bile duct proliferation with fibrosis | 92 (54) | 8 (5) | - | - | *** (15.6) |
| Kidney (n = 57) | | | | | |
| Glomerular diffuse capillary wall thickening | 72 (41) | 25 (14) | 3 (2) | - | *** (8) |
| Glomerular mesangial deposits/sclerosis | 22 (13) | 39 (22) | 39 (22) | - | * (5.6) |
| Interstitial fibrosis | 75 (42) | 12 (7) | 13 (8) | - | *** (49.8) |
| Tubular epithelial cell hyperplasia | 75 (42) | 12 (7) | 13 (8) | - | *** (6.7) |
| Tubular hyalinization/atrophy/dilatation/necrosis | 61 (35) | 12 (7) | 27 (15) | - | *** (30.8) |
| Tubular medullar hyaline casts | 84 (48) | 13 (8) | 3 (1) | - | * (6.6) |
| Mononuclear cell infiltrations | 39 (22) | 42 (24) | 19 (11) | - | n.s. |

Lesions are divided into groups of absent, mild, moderate and severe. n.s.: individuals with lesions not significantly older (mean age) than individuals without lesions at p > 0.05. *: individuals with lesions significantly older (mean age) than individuals without lesions at p ≤ 0.05, **: at p ≤ 0.01 and ***: at p ≤ 0.001.
Finally, the relationship between mercury concentration and each histologic change (absent vs. present) was tested by an age correcting analysis of covariance (Table 3). This was conducted with mercury concentration as dependent variable, age as covariable and histologic change (absent vs. present) as class variable, including their 1st order interaction links (age × histologic change). The statistical analyses were employed on all pooled individuals and additionally on subadults and adults, respectively. After a successive reduction of non-significant interactions, judged from the type-III sum of squares (p ≤ 0.05), the significance of each of the remaining factors was evaluated from the final model Least Square Mean (LSMean).

**Results**

**Liver lesions**

Multifocal mononuclear cell infiltrations (lymphocytes, and macrophages and neutrophils) were found in 11%, portal mononuclear cell infiltrations in 15% and lipid granulomas in 63% of the bears, respectively (Table 1). All animals showed hepatocytic foamy cytoplasm (microvesicular steatosis), while 64% presented lipid accumulating Ito cells and 76% exhibited macrovesicular steatosis (lipid vacuoles) in the periacinar zone 2 and 3. Mild bile duct proliferation accompanied by portal fibrosis was found in 8% of the animals. Only Ito cell lipid accumulation and bile duct proliferation were associated with age (both: p < 0.01) while none of the lesions were related to sex or season (all: p > 0.05).

**Renal lesions**

Glomerular diffuse capillary wall thickening (similar to membranous glomerulonephritis due to immune deposits on the epithelial side of the glomerular capillary base-ment membrane) and PAS positive glomerular mesangial deposits/fibrosis was found in 28% and 78% of the animals, respectively (Table 1). Interstitial fibrosis, including dense Masson Trichrome and PAS-positive total fibrous obliteration of the glomerular sclerosis, was found in 25% of the individuals. All glomerular lesions exhibited positive age relationships (p < 0.05). Hyperplasia of distal convoluted tubule and collecting duct epithelial cells was found in 25% of the bears. PAS-positive hyalinization of the proximal convoluted tubular basement membrane accompanied by proximal convoluted tubular dilatation, atrophy, necrosis and interstitial fibrosis was found in 39% of the animals. In moderate cases, these lesions were accompanied by interstitial fibrosis and total glomerular obliteration. Tubular cylindrical hyaline casts (protein) were found in the medulla of 16% of the individuals, indicating protein loss. All tubular lesions exhibited positive age relationships (p < 0.05). The prevalence of mild and moderate interstitial fibrosis was 25%, while mononuclear cell infiltration in cortex, medulla and papilla was recorded in 61% of the polar bears. Mononuclear cell infiltrations were not related to age (p > 0.05). None of the renal lesions were related to sex or season (summer vs. winter) (all: p > 0.05).

**Mercury concentrations**

Liver mercury levels ranged from 1.1–35.6 µg/g w.w. and renal levels ranged from 1–50 µg/g w.w., of which 2 liver values and 9 kidney values were above the known threshold toxic levels of 30 µg/g w.w. in terrestrial mammals (Table 2, Figure 1, 2). Mercury in subadults was significantly lower when compared to both males and females (p < 0.05), while kidney mercury levels in adult males were significantly higher when compared to adult females.

### Table 2: Mercury concentrations [mean ± SD (n), µg/g w.w.] in East Greenland polar bears

| Organ        | Subadults          | Adult females     | Adult males       |
|--------------|---------------------|-------------------|-------------------|
| Liver        | 6.27 ± 2.83 (32)*#  | 15.78 ± 8.6 (15)* | 14.99 ± 8.96 (12)*|
| Kidney       | 7.18 ± 3.32 (31)**##| 18.04 ± 10.97 (15)*| 29.42 ± 14.09 (11)**##|

*: Significantly higher when compared to subadults at p ≤ 0.05. **: Significantly higher when compared to adult females at p < 0.05. #: significantly positive relationship to age at p ≤ 0.002 (R² = 0.27). **##: significantly positive relationship to age at p ≤ 0.001 (R² = 0.45).

### Table 3: Significant results when testing histopathology vs. mercury levels in East Greenland polar bears

| Age Group | Organ        | Histopathology                  | p (n, F, R²) |
|-----------|--------------|--------------------------------|--------------|
| All       | Liver        | Ito cell*                       | 0.01 (50; 6.4; 0.53) |
| Adults    | Liver        | Lipid granulomas*               | 0.07 (22; 3.6; 0.38) |
| All       | Liver        | Portal bile duct proliferation and fibrosis*## | 0.007 (50; 8; 0.47) |
| Subadults | Kidney       | Tubular hyalinisation, atrophy, dilatation and necrosis*## | 0.07 (31; 3.4; 0.51) |

The calculations of differences between individuals with and without lesions are based on ANCOVA Least Square Mean regression analyses normalising for age. *: individuals with lesions higher in mercury levels than individuals without lesions. *##: individuals with lesions lower in mercury levels than individuals without lesions.
(p < 0.05). In general, the deviation was most pronounced in adult females. When analysing mercury vs. age relationship within subadults, adult females and adult males, respectively, the liver and kidney mercury increased significantly with age in subadults (both: p < 0.002), while that was not the case for adults. The analyses of covariance normalising for age revealed no difference in mercury concentrations when comparing all females and all males for both kidney (p > 0.05) and liver (p > 0.05) tissue, although it seemed that kidney concentrations decreased in old females and increased in old males. Neither was there a seasonal difference (summer vs. winter) in liver and kidney mercury concentrations (both: p > 0.05).

**Mercury concentrations and histological lesions**

Based on these results, the relationship between histopathology (present vs. absent) and mercury was analysed on the entire pooled material and in subadults and adults, separately, within the age-correcting analyses of covariance. The analyses showed that mercury liver levels were significantly higher in individuals exhibiting visible Ito cells when compared to those not exhibiting visible Ito cells (p < 0.02) (Table 3, Figure 3). A similar trend was found for liver lipid granulomas (p > 0.05) (Table 3, Figure 3). In case of portal fibrosis, mercury levels were significantly lower in individuals exhibiting portal fibrosis (p = 0.007) and the same trend was found in case of hyalinisation of renal tubular basement membranes (p = 0.07) (Table 3, Figure 3).

**Discussion**

**Liver histology**

We found several different histopathological changes that could not be ascribed to specific etiological factors. The mononuclear cell infiltrates were non-specific inflammatory reactions towards micro-organisms and/or injury of local blood vessels [32,33]. The bile duct proliferation and fibrosis were probably non-specific tissue reactions towards infections and thereby a result of ageing [32,33]. Prunescu et al. [34] speculated whether interstitial fibrosis was due to pre-hibernation physiological adaptations in brown bear (U. arctos). However, such a seasonal pattern could not be detected in the East Greenland polar bears. The number of lipid granulomas, in turn, was probably originated from Ito cell rupture. The lipid accumulation in polar bear Ito cells has previously been described and it constitutes the major accumulation and storage sites for lipophilic vitamin A [24,35-37]. Young bears only gradually start eating vitamin A rich prey until they are weaned at app. 2 years of age, which could explain the age difference in Ito cell numbers [36,37]. Specifically, the zonary (periacinarian) hepatocytic steatosis could be ascribed to high lipid ingestion and starvation while we cannot rule out other co-factors such as abnormal hepatocytic function and decreased synthesis of apo-proteins [32,33].

However, the considerable mercury concentrations accumulated in the liver tissue of the polar bears were in the range of adverse toxic effect levels for terrestrial mammals [1,2]. Specifically, hepatocytic steatosis was similar to mercury intoxication (hypoxia) in general [32,33]. The
Figure 2
Kidney levels of mercury (µg/g w.w.) in 32 male (left) and 25 female (right) East Greenland polar bears. Known threshold level for mercury toxicity in wildlife is given based on Dietz et al., (1998a) and AMAP (2005).

Figure 3
Mercury concentration vs. age divided on histological lesions (yes: square; no: triangle). Visible Ito cells = IC; portal bile duct proliferation and fibrosis = PB/F; lipid granulomas = LG and proximal convoluted tubular hyalinisation = TH.
signs of chronic inflammation, bile duct proliferation, portal fibrosis and hepatocytic steatosis were similar to those reported in mercury exposed bottlenose dolphins [10] and Arctic beluga whales [11]. Studies on laboratory rodents support these findings [38]. Furthermore, Dietz et al. [23] showed that as much as 94% of the mercury in East Greenland polar bear hair sampled during 1892–2001 was likely of anthropogenic origin. Such an increase demands certain tolerance from polar bears to avoid sub cellular organ damage via e.g. up regulated methallothionein synthesis and selenid complex binding [1,2,10,40]. Previous studies of East Greenland polar bears have shown that the liver mercury:selenium molar ratio exceeded 1:1 for a few individuals which indicate that some of the total mercury is on an ion toxic form [22]. The fact that portal bile duct proliferation and fibrosis was associated with decreasing mercury concentrations may be due to decreased liver metabolism/function (uptake) or a simultaneously hepatocytic subcellular injury or death [39,41,42]. Furthermore, the positive association between mercury concentration and number of Ito cells and lipid granulomas could indicate a direct impact from mercury on the prevalence of liver inflammation [24]. In conclusion, the liver lesions were a result of recurrent infections and age while mercury could not be ruled out as potential co-factor.

Renal histology
Diffuse thickening of the glomerular capillary wall, glomerular deposits/sclerosis and tubular changes are all well described in domestic animals [18,19], wildlife [11,17,40] and humans [15,16]. These lesions are all associated with age and recurrent infections while exposure to toxic substances is a known co-factor [15,16]. In the polar bears, glomerular sclerosis was worsen in cases of interstitial fibrosis, which is an expected and age-related change in various mammals, while the observed tubular epithelial cell hyperplasia has been associated with regeneration of renal parenchyma and chronic renal failure [15,16]. The interstitial nephritis (mononuclear cell infiltrations) was similar to those found in seals from North West Greenland and the Baltic [17,40] which we ascribe to chronic recurrent infections [15,16,18,19].

However, the histopathological lesions found in glomeruli (immune-complex glomerulonephritis), tubules and interstitium resemble those in Baltic grey seal and ringed seal heavily polluted with heavy metals between 1977–1996 [17]. Similar associations are also reported from controlled laboratory studies [12-16], in humans [15,16], in domestic mammals [18,19] and other Arctic wildlife species [11]. Proximal convoluted tubular basement membrane hyalinisation was associated with decreasing mercury levels. That could be due to a decreased metabolism/kidney function or because of mercury induced liver injury. Regarding the latter possibility, the quoted hepatic damage would increase the leakage of mercury (included in polypeptide complexes) into plasma and hence, it would also increase the burden of Hg filtered through the glomeruli, which in turn would generate tubular hyalinisation [39,41,42] Proximal convoluted tubular basement membrane hyalinisation was associated with decreasing mercury levels. That could be due to a decreased metabolism/kidney function or because of mercury induced liver injury. Regarding the latter possibility, the quoted hepatic damage would increase the leakage of mercury (included in polypeptide complexes) into plasma and hence, it would also increase the burden of Hg filtered through the glomeruli, which in turn would generate tubular hyalinisation [39,41,42]. According to Dietz et al. [22], the East Greenland polar bear renal tissue mercury:selenium molar ratio exceeded 1:1 for several individuals, which indicate that a large amount of the total mercury is on the ion toxic form, suggesting a toxic potential. However, the conclusion is that the chronic nature of the lesions was a result of age and recurrent infections while we cannot exclude mercury as a co-factor.

Considerations
Whether the present lesions have an impact on health status at the individual level is impossible to evaluate. However, it cannot be excluded that individuals that are more susceptible to mercury toxicity may be affected as 16% and 3% of the animals exceeded renal and liver threshold levels, respectively, for toxic effects of the quoted metal in mammals [2]. Furthermore, Sonne et al. [24,25] showed that anthropogenic organochlorines and brominated flame retardants were possible co-factors in the development of renal and liver lesions in East Greenland polar bears. The present study shows associations between organ lesions and mercury levels which indicate mercury as a co-factor although recurrent infections and age clearly were main factors. Anyhow, the mercury levels adds another risk factor to polar bear health as the pollution cocktail increase and thereby the possibility of additive effects. Furthermore, the polar bear health mercury problem may be most pronounced in the Western Arctic, as the highest concentrations exists there [1,2,23] due to increasing emissions from Eurasia.

Conclusion
We found liver and kidney mercury levels in East Greenland polar bears in the range of toxic effects. The signs and nature of chronic inflammation and statistical relationships points towards age and recurrent infections as main factor while mercury could not be ruled out as a possible co-factor. These are new and important results in the monitoring and assessment of the potential toxic impact from the increasing mercury concentrations in Arctic wildlife and humans relying on polluted marine species.
Competing interests
The author(s) declare that they have no competing interests.

Authors’ contributions
CS drafted the manuscript. CS and RDJ conducted the age determinations with MKI. PSL participated in the histopathological examinations. GA conducted the mercury analyses, EWB participated in the study design. All authors read and approved the final manuscript.

Acknowledgements
Danish Cooperation for Environment in the Arctic for financial support, H. & B. Sandell, Jonas Brønlund and local hunters for organizing sampling in East Greenland. Sigga E. Joensen and Lene Bruun for conducting mercury analyses. Two reviewers are acknowledged for their comments on the submited manuscript.

References
1. Arctic Monitoring and Assessment Programme: AMAP Assessment Report – Arctic Pollution Issues. Oslo 1998 [http://www.amap.no]
2. Arctic Monitoring and Assessment Programme: AMAP Assessment 2002 – Heavy Metals in the Arctic. Oslo 2005 [http://www.amap.no]
3. Berg T, Bartnicki J, Munthe J, Lattila H, Hrehoruk J, Mazur A: Atmospheric mercury species in the European Arctic: Measurements and modelling. Atmos Environ 2001, 35:2569-2582.
4. Lindberg SE, Brooks S, Lin CJ, Scott K, Meyers T, Chambers L, Landis M, Stevens R: Formation of reactive gaseous mercury in the Arctic: evidence of oxidation of Hg0 to gas-phase Hg-II compounds after Arctic sunrise. Water Air Soil Pollut 2001:295-302.
5. Lu JY, Schroeder WH, Barrie L, Steffen A, Welch H, Martin K, Lockhart L, Hunt R, Boila G, Richter A: Magnification of atmospheric mercury deposition to polar regions in springtime: the link to tropospheric ozone depletion chemistry. Geophys Res Lett 2001, 28:3219-3222.
6. Schroeder WH, Anlauf KG, Barrie LA, Lu JY, Steffen A, Schneeberger DR, Berg T: Arctic springtime depletion of mercury. Nature 1998, 394:331-332.
7. Arctic Monitoring and Assessment Programme: AMAP Assessment 2002 – Human Health in the Arctic. Oslo 2003 [http://www.amap.no]
8. Goyer RA, Clarkson TW: Toxic effects of metals. In Cassaret and Doul’s Toxicology: The Basic Science of Poisons Edited by: Klaassen CD. New York: McGraw-Hill; 2001:811-868.
9. Hylander LD, Goodsite M: Environmental costs of mercury pollution. Sci Total Environ 2006, 368:332-370.
10. Rawson AJ, Patton GW, Hofmann S, Pietra GG, Johns L: Liver abnormalities associated with chronic mercury accumulation in stranded Atlantic bottlenose dolphins. Ecotoxicol Environ Saf 1993, 25:41-47.
11. Wohrer VM, O’Hara TM, Eurell JA, Willig MA, Bratton GR, Suydam JR, Beasley VR: Inorganic mercury in liver and kidney of beluga and bowhead whales through autometallo genesis. J Pathol 1991, 164:407-408.
12. Afonin OJ, Orisakwe OE, Obi E, Dioksa CE, Ndubuka GI: Nephro toxic actions of low-dose mercury in mice: protection by zinc. Arch Environ Health 2002, 57:9-102.
13. Banerjee S, Bhattacharya S: Histopathology of kidney of channa punctatus exposed to chronic nonlethal level of elsan, mercury and ammonia. Ecotoxicol Environ Safety 1994, 29:265-275.
14. Rumbeha WK, Fitzgerald SD, Braselton WE, Roh TA, Pesta JK, Kanene JB: Augmentation of mercury-induced nephrotoxicity by endotoxin in the mouse. Toxicology 2000, 151:103-116.
15. Chung J, Bernasek J, Glassock RJ: Renal disease, Classification and atlas of glomerular diseases. New York: Igaku-Shoin; 1995.
16. Cotran RS, Kumar V, Collins T: Robbins pathologic basis of disease. Philadelphia: WB Saunders Company; 1999.
17. Bergman A, Bergstrand B, Bignert A: Renal lesions in Baltic grey seals (Halichoerus grypus) and ringed seals (Phoca hispida botnica). Ambio 2001, 30:397-409.
18. Confer AW, Panciera RJ: Thomsom’s Special Veterinary Pathology. St. Louis: Mosby Year Book; 1995.
19. Maxie MG: Pathology of domestic animals. San Diego: Academic Press; 1993.
20. Ramsay MA, Stirling I: Reproductive biology and ecology of female polar bears (Ursus maritimus). J Zool 1988, 214:601-634.
21. Dietz R, Overgaard Nielsen C, Munk Hansen M, Hansen CT: Organic mercury in Greenland birds and mammals. Sci Total Environ 1990, 95:41-51.
22. Dietz R, Rigit FF, Born EW: Geographical differences of zinc, cadmium, mercury and selenium in polar bears (Ursus maritimus) from Greenland. Sci Total Environ 2000, 245:25-48.
23. Dietz R, Rigit FF, Born EW, Kirkegaard M, Olsen MT, Asmund G, Renzonni A, Baagøe H, Andresen C: Trends in Mercury in Hair of Greenlandic Polar Bears (Ursus maritimus) during 1892–2001. Environ Sci Technol 2006, 40:1120-1125.
24. Sonne C, Dietz R, Leifsson PS, Born EW, Kirkegaard M, Rigit FF, Lechtrek RJ, Muir D GG, Hyldstrup L: Are organohalogen contaminants a co-factor in the development of renal lesions in East Greenland polar bears (Ursus maritimus)? Environ Toxicol Chem 2002, 25:1551-1557.
25. Dietz R, Heide-Jorgensen MP, Harkønen T, Teilmann J, Valentin N: Age determination of European harbour seal (Phoca vitulina L.). J. Zool 1991, 176:17-21.
26. Rosing-Asvid A, Born EW, Kingsley MCS: Age at sexual maturity of males and timing of the mating season of polar bears (Ursus maritimus) in Greenland. Polar Biol 2001, 25:876-883.
27. Derchoer AE, Stirling I: Age-specific reproductive-performance of female polar bears (Ursus maritimus). Can J Zool 1994, 34:527-536.
28. Bancroft JD, Stevens A: Theory and Practice of Histological Techniques. New York: Churchill Livingstone; 1995.
29. Lyon H, Andersen AP, Hasselager E, Hayer PE, Møller M, Prenta Van Deurs B: Theory and Strategy in Histochemistry. Berlin: Springer Verlag; 1991.
30. Asmund G, Vorkamp K, Backus S, Comba M: An update on analytical methods, quality assurance and quality control used in the Greenland AMAP programme 1999 to 2002. Sci Total Environ 2004, 331:233-245.
31. Kelly WR: The liver and biliary system. In Pathology of Domestic Animals Edited by: Jubb KVF, Kennedy PC, Palmer N. San Diego: Academic Press; 1993:319-406.
32. MacLachlan NJ, Cullen JM: Liver, biliary system and exocrine pancreas. In Thomsom’s Special Veterinary Pathology Edited by: Carlton WW, Donald McGavin M. St. Louis: Mosby Year Book; 1995:81-115.
33. Prunescu CC, Serban-Parau N, Brock JH, Vaughan DM, Prunescu P: Liver and kidney structure and iron content in romanian brownbears (Ursus arctos) before and after hibernation. Comp Biochem Physiol A 2003, 134:1-26.
34. Leighton FA, Cattet M, Norstrum R, Trudue S: A cellular basis for high-levels of vitamin-A in livers of polar bears (Ursus maritimus) – the Ito cell. Can J Zool 1994, 72:480-482.
35. Senoo H, Imai K, Wake K, Wold H, Moskagj K, Kojima N, Matano Y, Miura M, Sato M, Roos N, Berg T, Langvatn R, Norum K, Blomhoff R: Vitamin A-storing system in mammals and birds in an Arctic area – a study in the Svalbard Archipelago. In Cells of the hepatic sinusoid Edited by: Wiss E, Knook DL, de Zanger R, Fraaser R, Leiden: Kupffer Cell Foundation; 1999:34-35.
36. Senoo H, Imai K, Higashi N, Wake K, Kojima N, Miura M M, Wold H, Moskagj K, Sato T, Sato M, Roos N, Berg T, Norum K, Blomhoff R: Transport and hepatic storage of vitamin A in arctic animals. In Cells of the hepatic sinusoid Edited by: Wiss E, Knook DL, de Zanger R, Fraaser R, Leiden: Kupffer Cell Foundation; 2001:207-209.
37. Al-Saleh I, Khogali F, Al-Amodi M, El-Dousch I, Shinwari N, Al-Baradei R: Histopathological effects of mercury in skin-lightening cream. Environ Pathol Toxicol Oncal 2003, 22:287-299.
39. Dietz R, Paludan-Müller P, Thye Agger C, Nielsen CO: Cadmium, mercury, zinc and selenium in ringed seals (Phoca hispida) from Greenland waters. Nammco Scientific Contributions 1998, 1:242-273.

40. Sonne-Hansen C, Dietz R, Leifsson PS, Hylistrup L, Riget FF: Cadmium toxicity to ringed seals (Phoca hispida) – An epidemiological study of possible cadmium induced nephropathy and osteodystrophy in ringed seals (Phoca hispida) from Qaanaaq in Northwest Greenland. Sci Total Environ 2002, 295:67-181.

41. Dietz R, Riget FF, Johansen P. Lead, cadmium, mercury and selenium in Greenland marine animals. Sci Total Environ 1996, 186:67-93.

42. Dietz R, Nørgaard J, Hansen JC. Have arctic marine mammals adapted to high cadmium levels? Mar Pollut Bull 1998, 36:490-492.