We analyzed bone mineral density (BMD) in skulls of polar bears (*Ursus maritimus*) (*n* = 139) from East Greenland sampled during 1982–2002. Our primary goal was to detect possible changes in bone mineral content (osteopenia) due to elevated exposure to organochlorine (polychlorinated biphenyls (PCBs), dichlorodiphenyl trichloroethane (DDT), and its metabolites, chlordanes (CHLs), dieldrin, hexachlorocyclohexanes, hexachlorobenzene) and polybrominated diphenyl ether (PBDE) compounds. To ensure that the BMD value in skull represented the mineral status of the skeletal system in general, we compared BMD values in femur and three lumbar vertebrae with skull in a subsample. We detected highly significant correlations between BMD in skull and femur (*r* = 0.99; *p* < 0.001; *n* = 13) and skull and vertebrae (*r* = 0.97; *p* < 0.001; *n* = 8). BMD in skulls sampled in the supposed pre-organochlorine/PBDE period (1892–1932) was significantly higher than in skulls sampled in the supposed pollution period (1966–2002) for subadult females, subadult males, and adult males (all, *p* < 0.05) but not adult females (*p* > 0.94). We found a negative correlation between organochlorines and skull BMD for the sum of PCBs (ΣPCB: *p* < 0.04) and ΣCHL (p < 0.03) in subadults and for dieldrin (p < 0.002) and ΣDDT (p < 0.02) in adult males; indications for ΣPBDE in subadults were also found (*p* = 0.06). In conclusion, the strong correlation relationships suggest that disruption of the bone mineral composition in East Greenland polar bears may have been caused by organochlorine exposure. Key words: BMD, bone mineral density, chlordane, DDT, dieldrin, endocrine disruption, osteoporosis, PCBs, polar bear, polychlorinated biphenyls, *Ursus maritimus*. Environ Health Perspect 112:1711–1716 (2004). doi:10.1289/ehp.7293 available via http://dx.doi.org/ [Online 13 September 2004]
up-concentrate in the blubber of ringed seal (*P. hispida*) and bearded seal (*Etrangisatus barbatus*), which is the primary food of the polar bear (AMAP 2004; de March et al. 1998; Lie et al. 2003; Norstrom et al. 1998). Recent studies of polar bears from Svalbard have indicated that high levels of organochlorines negatively affect levels of retinol (vitamin A) and thyroid hormones (Bræathen et al. 2004; Skaare et al. 2001) and possibly also negatively affect cortisols, sex steroids, and reproductive organs (female pseudohermaphrodites), although these latter mechanisms are not clearly understood (Haave et al. 2003; Oskam et al. 2003, 2004; Sonne et al., in press; Wig et al. 1998). Other studies have associated high levels of organochlorines with low levels of IgG, suggesting possible immunotoxic effects on the IgG levels (Bernhoft et al. 2000; Lie E, Larsen HJS, Larsen S, Johansen GM, Derocher AE, Lunn NJ, et al., unpublished data). Overall, these studies support the notion that organochlorines may cause disruption and thereby potentially affect bone mineral composition.

To determine whether exposure to organochlorines and PBDEs may have adversely affected bone mineral composition in polar bears, we compared BMD in skulls of 41 individual polar bears collected in East Greenland during the supposed prepolluted period (1892–1932) with 98 polar bear skulls collected during the supposed polluted period (1966–2002). Furthermore, we examined a subset of 58 of the individuals collected during the pollution period to determine if BMD was related to body burden of various organochlorines and PBDEs.

**Materials and Methods**

**Sampling and age estimation.** We studied a total of 139 East Greenland polar bear skulls sampled between Skjoldungen at 63°15’N and Danmarks Havn at 76°30’N during 1892–2002. The age determination was carried out by counting the cementum growth layer groups (GLGs) of the lower left incisor (I3) after decalcification, thin sectioning (14 µm), and staining (toluidine blue) using the method described by, for example, Hensel and Sorensen (1980) and Dietz et al. (1991). For analyses, the individuals were then categorized into adult males (≥ 6 years of age), adult females (≥ 5 years), and subadults (others) (e.g., Rosing-Asvid et al. 2002). Regarding skull samples from 1892–1987, the sex was available from the expedition files, and in case of absence of this information (n = 9), the determination was based on skull morphology.

**Osteodensitometry.** X-Ray osteodensitometry was applied to detect osteopenia (osteoporosis) by use of an X-ray bone densitometer (model XR 26; Norland Corporation, Fort Atkinson, WI, USA), which determined the BMD (calcium phosphate, hydroxyapatite) using dual X-ray absorptiometry (DXA). The skulls were scanned in “research” mode (speed, 60 mm/sec; resolution, 3.0 × 3.0 mm; width, 24.9 cm) and analyzed using XR software (revision 2.4; Norland Corporation), which generated a picture of the bone segment and calculated the BMD of hydroxyapatite in grams per square centimeter (Figure 1).

To ensure that BMD in the skull represents the mineral status of the skeletal system in general, a study was conducted where the BMD of the skull, one femur, and three lumbar vertebras were compared in a subset of 13 free-ranging polar bears (3 subadults, 2 adult females, and 8 adult males) from Svalbard and East Greenland. The DXA scanner was calibrated daily using a phantom with known mineral density. In addition, the precision was tested by a 10× rescanning (mean ± SD, 521.96 ± 0.60 g/cm²), which from the formula [1 – (SD/mean) × 100%] gives a precision of 99.88%. Fragmentation and loss of tooth material caused by handling and lead shot were thought to be a problem. A double determination of the BMD in 2 skulls (numbers 5483 and 2891) with and without incisors, canines, premolars, and molars showed that loss of half or more of the material of the large canines altered the result significantly. Because the canines in the material were not fragmented to such a degree, we did not consider fragmentation a problem.

**Contaminant analyses.** Polar bear subcutaneous adipose tissue samples (n = 58) were analyzed for PCBs, DDT and its metabolites, HCHs, CHLs, HCB, dieldrin, and PBDEs as described elsewhere (Dietz et al. 2004; Luoss et al. 2002; Sandala et al. 2004). The sum of PCBs (ΣPCB) is the total concentrations of the 51 individual or coeluting congeners (if detected); PCBs 31/28, 52, 49, 44, 42, 64/71, 74, 70, 66/95, 60, 101/84, 99, 97, 87, 110, 151, 149, 118, 146, 153, 105, 141, 179, 138, 158, 129/178, 182/187, 183, 128, 174, 177, 171/202/156, 200, 172, 180, 170/190, 201, 203/196, 195, 194, and 206. ΣDDT is the sum of 4,4′-DDT, 4,4′-DDD (dichlordiphenyldichloroethane), and 4,4′-DDE (dichlorodiphenyldichloroethylene). ΣHCH is the sum of the α-, β-, and γ-hexachlorocyclohexane. ΣCHL is the total concentration of oxychlordane, trans-chlordane, nonachlor III (MC6), trans-nonachlor, cis-nonachlor, and heptachlor epoxide. ΣPBDE concentration is the total of 35 individual or coeluting congeners (if detected); PBDE numbers 10, 7, 11, 8, 12/13, 15, 30, 32, 28/33, 35, 37, 75, 71, 66, 47, 49, 77, 100, 119, 99, 116, 85, 155/126, 105, 154, 140, 138, 166, 183, 181, and 190 (Muir DCG, Dietz R, Riget FF, Sonne C, Letcher RJ, Born EWB, unpublished data). All contaminant data are given in nanograms per gram lipid weight (l.w.).

**Statistics.** The BMD showed no deviation from normality (Shapiro–Wilks test), whereas contaminant data were log-transformed (base e) before analyses in order to meet the criteria of normality and homogeneity of the variance. The significance level was set to p ≤ 0.05, and a significance level of 0.05 < p ≤ 0.10 was considered a trend. First, we tested the condylar skull length versus age within each group (i.e., subadults of both sexes, adult females, and adult males) in an analysis of covariance (ANCOVA) with skull length as a dependent variable, periods (before and after 1960 respectively) as class variables, age as a covariable, and their first-order interaction links (age × period). The result from this analysis showed that the relationship of skull length versus age was the same in the two periods, which justified the use of non-length-corrected skull BMD in the further analyses (all, p > 0.26). Second, the relationship of BMD versus age was tested by a linear regression analyses (BMD as a dependent variable and age as an independent variable) for subadults of both sexes, adult females, and adult males. To test for period differences, we used an ANCOVA with BMD as a dependent variable, age/sex (subadult females, subadult males, adult females, and adult males) and period (before and after 1960 respectively) as class variables, age as a covariable, and the first-order interaction links (age × period, age × age/sex, and age/sex × period) between these variables.

![Figure 1](image-url)
variables. The model was successively reduced for nonsignificant interactions (p > 0.05) judged from the type III sum of squares, and the significance of the remaining factors was evaluated from the final model (least square means). A temporal trend over the entire period 1892–2002 was analyzed by a multiple regression analysis with skull BMD as the dependent variable and the individual age and year of kill as explanatory variables for subadults of both sexes, adult females, and adult males, respectively (the relationship was evaluated from the parameter estimate, r², and p-value). The relationship between age/sex groups and contaminants was analyzed within a one-way analysis of variance on the log-transformed contaminant data, and significant results were tested by Tukey’s post hoc test.

The skull BMD versus contaminant (ΣPCB, ΣDDT, ΣCHL, HCB, ΣCHD, dieldrin, and ΣPBDE) relationships were explored by multiple regressions with skull BMD as the dependent variable and the age and contaminant concentrations as explanatory variables within age/sex groups (subadults of both sexes, adult females, and adult males). Finally, the relationship between levels of contaminants and BMD was evaluated from the parameter estimate, r², and p-value.

Results

We found a highly significant correlation between BMD in skull and femur (r = 0.99; p < 0.001; n = 13), and skull and vertebrae (r = 0.97; p < 0.001; n = 8). These results justified the use of BMD measurements in skull to reflect the status of the skeletal system, although information on body conditions and nutritional stressors, relevant for osteoblastic and osteoclastic activity, was not available.

Skull BMD and age/sex differences. BMD was analyzed in 139 skulls representing the period 1892–2002, and consisted of 64 subadults, 40 adult females and 35 adult males. The BMD increased with age in subadults (p < 0.001) but not adults (both, p > 0.05; Figure 2). BMD differed between males and females (p < 0.01) in the order subadult females < subadult males < adult females < adult males. Furthermore, BMD in females 14–23 years of age seemed to decline significantly with age (p < 0.04).

Discussion

BMD and age/sex differences. The high correlation in BMD between skull and femur and vertebrae, respectively, is useful because skull samples of polar bears (and other mammals) are present at national zoological museums all over the world, which makes various time-trend bone studies possible. Our results clearly show that skull BMD increased more rapidly in subadults compared with adults, in accordance with previous studies of ringed seals from Northwest Greenland (Sonne-Hansen et al. 2002). Female polar bears usually give birth to two cubs every third year (December) and mobilize and transfer large amounts of calcium and phosphate during gestation and during the postpartum (suckling) period, which lasts up to 2 years (Ramsay and Stirling 1988). In this period, calcium is used for fetal skeletal production and maintenance of the mother’s and offspring’s calcium phosphate homeostasis (Ramsay and Stirling 1988). Because the female polar bear mobilizes these large amounts of calcium and phosphate, adult females are expected to have a lower BMD compared with adult males. Such a difference was also found in the present study. Similar differences have been found in humans (e.g., Van Langendonck et al. 2002). As suggested for humans, the marked difference in BMD between the sexes could be the result of a higher muscle mass and strength in males, leading to higher biomechanical loading of the bone. This would lead to increased bone formation through the stimulation of the mechanotransduction system in the osteocytes (Van Langendonck et al. 2002).

Earlier studies show that sufficient levels of sex steroids (estrogens and androgens) are important in the development of the human cortical bone structures in boys, girls,
teenagers, adults, and the elderly (Hampson et al. 2002; Juul 2001; Leder et al. 2001; Szulc et al. 2001). On the other hand, high levels of estrogen-active substances (intrinsic, extrinsic) stimulate the expression of secondary sexual characteristics (Hampson et al. 2002; Juul 2001; Leder et al. 2001; Szulc et al. 2001). Therefore, growth delay and osteopenia (osteoporosis) have been associated with hypogonadism and lower estrogen levels in both sexes (Leder et al. 2001; Nelson 2003; Szulc et al. 2001). The age-related decrease in BMD in females in the present study was probably associated with a menopause phase after 15 years of age, but this requires a larger sample size (Figure 2) (Derocher and Stirling 1994).

**Period differences and temporal trends in skull BMD.** In both analyses of subadults of both sexes and adult males, the individuals from the prepollution period had a higher skull BMD compared with those from the polluted period. These results suggest that there is a linkage between decreased BMD for bears from the polluted period and exposure to environmental stressors compared with bears in the prepollution period. Two major environmental stressors could be linked to mineral loss in polar bear skulls: anthropogenic organochlorine compounds and PBDEs and/or climate oscillations (AMAP 2004; de March et al. 1998; Farland et al. 2002). Concentrations of, for example, ΣPCB in the adipose tissue of East Greenland polar bears have, over the last four decades, reached levels that can elicit adverse biological effects on immunologic parameters and vitamin A levels, which may be linked to the present decrease in skull BMD (stress) (AMAP 2004; de March et al. 1998). However, during the same period global warming has resulted in a reduction in the ice coverage in the East Greenland area (Comiso 2002; Rothrock et al. 1999). Although population ecology has not been studied in East Greenland, the situation is probably similar for polar bears from the Hudson Bay area in Canada (Stirling et al. 1999). A reduction of the sea ice in the Hudson Bay area has reduced the bears’ access to ringed seals, resulting in reduced body condition and lowered natality in the polar bears (Stirling et al. 1999).

**Temporal differences with respect to potential effects of PCB and DDT exposure on periodontitis and osteoporosis in gray seal and harbor seal was investigated by Bergman et al. (1992), Mortensen et al. (1992), and Schandorff (1997). They found exostosis and periodontitis, often with substantial loss of alveolar bone in mandible and maxilla (osteoporosis). These changes could have been caused by hormonal imbalance potentially induced by PCBs and by DDT and its metabolites, with malformation of the calcium helix structures around the collagen matrix (DeLillis 1989). These results are further supported by the investigations of Render et al. (2000a, 2000b, 2001). However, it must be noted that the ranges of ΣPCB and ΣDDT levels in the seals were orders of magnitude higher compared with levels in the present polar bears (Blomkvist et al. 1992).**

Lind et al. (2003) investigated the BMD in the male gray seals (n = 43) reported by Bergman et al. (1992). The method used was peripheral quantitative computed tomography, which made it possible to distinguish between cortical and trabecular bone in os mandibularis and os radius, respectively (DXA scanning used in the present study gives the average of trabecular and cortical bone density). Three sample groups of seals were compared: 1850–1955 (no pollution), 1965–1985 (high pollution), and 1986–1997 (fairly low pollution). They found that trabecular BMD was significantly higher in the fairly low pollution period (1986–1997) compared with the high pollution period (1965–1985), whereas mandible cortical BMD was significantly lower in the fairly low pollution period (1986–1997) compared with the no-pollution period (1850–1955). Our study of BMD in East Greenland polar bear supports the findings of Lind et al. (2003).

**BMD levels and contaminants.** Bone density expresses the bone mineral composition determined by the activity of osteoblastic bone formation and osteoclastic bone resorption, which is regulated by androgens and estrogens through cytokines (Manolagas and Jilka 1995; Manolagas et al. 1995). Studies on Svalbard have shown that PCBs may negatively influence plasma cortisol, estrogen, and testosterone levels (Haave et al. 2003; Oskam et al. 2003, 2004) and plasma retinol and thyroid hormone levels in polar bears (Braathen et al. 2004; Skaare et al. 2001). These studies all indicate that organochlorines in Svalbard polar bears (and likely also East Greenland bears, because the organohalogen compound levels are comparable) potentially affect endocrine homeostasis, which again may lead to bone mineral loss (osteoporosis). Another polar bear study from Svalbard associated high levels of organochlorines with low levels of IgG, suggesting possible immunotoxic effects (Bernhoff et al. 2000; Lie E, Larsen HJS, Larsen S, Johansen GM, Derocher AE, Lunn NJ, et al., unpublished data). This potential effect may lower the immune response and enhance stress with increased cortisol levels, which potentially affects the bone mineral composition (osteoporosis).

The present study indicated that high concentrations of ΣPCB and ΣCHL are associated with reduced skull BMD in subadults and that ΣDDT and dieldrin are associated with reduced skull BMD in adult males. These BMD relationships with ΣPCB, ΣCHL, ΣDDT, and dieldrin concentrations in subadults of both sexes and adult males may suggest endocrine-related effects (e.g., AMP 2002; Birnbaum 1994; Damstra et al. 2002; de March et al. 1998; Lind et al. 2003, 2004). For example, PCBs and DDT and its metabolites have shown in vitro and in vivo to be weak agonists/antagonists of estrogen-receptor-mediated activity; organochlorine-

**Table 3. Concentrations [mean ± SD (median), ng/g l.w.] of various contaminants in subcutaneous adipose tissue of 58 East Greenland polar bears sampled during 1999–2001.**

| Compound | Subadults (n = 35) | Adult females (n = 14) | Adult males (n = 9) |
|----------|--------------------|------------------------|---------------------|
| ΣPCB     | 6.597 ± 2.726 (6.089) | 5.334 ± 2.150* (5.770) | 8.637 ± 4.111* (8.280) |
| ΣCHL     | 1.598 ± 0.884 (1.469) | 1.379 ± 0.591 (1.353) | 1.055 ± 0.517 (0.914) |
| ΣDDT     | 392 ± 209 (376)     | 358 ± 149 (366)      | 481 ± 331 (496)     |
| ΣHCH     | 196 ± 68 (172)      | 195 ± 106 (151)      | 294 ± 210 (181)    |
| Dieldrin | 210 ± 100 (186)     | 174 ± 70 (154)       | 177 ± 81 (172)     |
| HCB      | 99 ± 84 (70)        | 75 ± 82 (61)         | 51 ± 28 (48)       |
| ΣPBDE    | 62 ± 23 (53)        | 53 ± 17 (53)         | 52 ± 16 (49)       |

*Significant difference between adult females and males at the p ≤ 0.05 level.

**Table 4. Significant results from the multiple regression analyses of skull BMD versus age and contaminant concentrations in East Greenland polar bears sampled during 1999–2001.**

| Age/sex group | Equation | $r^2$ | $\rho_{\text{age}}$ | $\rho_{\text{cont}}$ | No. |
|---------------|----------|-------|----------------------|----------------------|-----|
| Subadults     | BMD = 0.26 × age – 0.25 × ln(ΣPCB) + 3.1 | 0.58  | < 0.001             | < 0.04**              | 35  |
| Adult males   | BMD = 0.24 × age – 0.19 × ln(ΣHCH) + 2.4  | 0.6   | < 0.001             | < 0.03**              | 35  |
|               | BMD = 0.25 × age – 0.18 × ln(ΣPBDE) + 1.69 | 0.58  | < 0.001             | 0.06*                 | 9   |
| Adult males   | BMD = 0.005 × age – 0.37 × ln(dieldrin) + 4.5 | 0.85  | 0.43                | 0.002*                | 9   |

The equation is given as [BMD = A × age + B × ln(contaminant) + C], with BMD (g/cm²) as the dependent variable and age (years) and log-transformed contaminant concentration (ln(ng/g l.w.) as the explanatory variables. A, B, and C are specific parameter estimates; $r^2$ is the regression coefficient of the model; $\rho_{\text{age}}$ is the p-value for age; and $\rho_{\text{cont}}$ is the p-value for contaminants.

*Nonsignificant trend of a negative correlation between BMD and ln(ΣPBDE) at the 0.05 < p ≤ 0.10 level. **Significant negative correlation between BMD and organochlorine contaminant group at the p ≤ 0.05 level. *Significant negative correlation between BMD and organochlorine contaminant group at the p ≤ 0.01 level.
mediated induction of cytochrome P450 isozyme activity can affect circulating sex hormone levels (e.g., estrogens) (Navas and Segner 1998), and this is also of relevance in the polar bear (e.g., Letcher et al. 1996). Relationships between 4,4’-DDE concentrations and BMD in humans have been reported (Beard et al. 2000; Glynne et al. 2000). Glynne et al. (2000) found significant negative correlations between 4,4’-DDE and BMD in 68 sedentary women (where concentrations are lower compared with the present polar bears) and concluded that 4,4’-DDE may also have a negative effect on BMD in men (with contaminant levels comparable with those found in the polar bears). Lind et al. (2004) investigated the relationship between DDT and its metabolites and bone composition in juvenile female American alligators (Alligator mississippiensis) in Lake Apopka, Florida. Compared with data from a nonpolluted reference alligator subpopulation, the trabecular bone mass was increased, and the authors suggested that environmental estrogens (e.g., DDT and its metabolites) disrupted the normal bone remodeling process (inhibition of osteoclast activity), which had resulted in increased BMD.

Guo et al. (1994) found that children (n = 25) of primiparous PCB-contaminated mothers (Yu-Cheng rice oil disease) were significantly smaller and had less total lean mass and less soft tissue mass but not lower BMD compared with a control group. The PCB levels in the children (serum) were 10.3 ng/g l.w., which is lower than the levels in polar bears in the present study. Alveblom et al. (2003) investigated the incidence of osteoporotic fractures in fishermen and their wives from the Baltic Sea (high pollution) and compared these with fishermen from the west coast of Sweden (low pollution) as controls. For vertebral fractures, there was a significantly higher incidence rate ratio for east coast (Baltic) women compared with west coast women, and a similar but nonsignificant tendency was found for men. The PCB concentration (10 congeners) was 2,000 ng/g l.w., which was significantly higher compared with the west coast population but lower compared with the range in the subsurface adipose tissue of East Greenland polar bears. These environmental studies support the findings of negative associations between PCBs/DDT and BMD levels in East Greenland polar bears.

In the present study, we observed a negative correlation between 2PBDE concentrations in adipose tissue and BMD in subadults. Disturbances in thyroid function and developmental toxicity (histopathology) have been shown to be associated with PBDEs in laboratory rats (e.g., de Wit 2002) as well as in polar bears from Svalbard (Braathen et al. 2004; Skaare et al. 2001).

Conclusions

Skull BMD increased with age in subadults and was higher in males than in females at all ages. For adult females from 14 years of age, a menopausal BMD decrease was indicated, but further examination requires a larger sample size. BMD in skulls from subadult females, subadult males, and adult males sampled in the supposed pollution period (1966–2002) was significantly lower than BMD in skulls from the period before supposed pollution with organochlorines and PBDE compounds (1892–1932). Furthermore, correlative relationships suggest that ΣPCB, ΣChl, dieldrin, and 2,3,7,8-∑DDT exposure negatively influenced BMD in skulls from both sexes and adult males.

Correction

In the manuscript originally published online, the years 1892–1960 and 1961–2002 were used to represent the pre- and post-organochlorine/PBDE periods, respectively. These years have been changed throughout to reflect the years in which the skulls were actually collected (1892–1932 and 1966–2002).

References

Alveblom AK, Rylander L, Johneil D, Hagmar L. 2003. Incidence of hospitalized osteoporotic fractures in cohorts with high dietary intake of persistent organic compounds. Int Arch Occup Environ Health 76:246–248.

AMA. 2004. Persistent Organic Pollutants in the Arctic. AMAP Assessment 2002. OsloArctic Monitoring and Assessment Programme.

Bead J, Marshall S, Jong K, Newton R, Tripplett-McBride T, Humphries E, et al. 2000. 1,1,1-Trichloro-2,2,2-bis (p-chlorophenyl)-ethane (DDT) and reduced bone mineral density. Arch Environ Health 55(3):177–180.

Bengtsson BE, Bengtsson A, Himberg M. 1996. Fish deformities and pollution in some Swedish waters. Am. 142:32–35.

Bergman A, Olsson M, Reiland S. 1992. Skull bone lesions in the Baltic grey seal (Halichoerus grypus). Diploma. 21:517–519.

Bernholt A, Skare JU, Wiig Ø, Derocher AE, Larsen NJU. 2000. Possible immunotoxic effects of organochlorines in polar bears (Ursus maritimus) at Svalbard. J Toxicol Environ Health A 57(7):561–574.

Bertholdt A, Wiig Ø, Skare JU. 1997. Organochlorines in polar bears (Ursus maritimus) at Svalbard. Environ Pollut 98:159–175.

Birnbaum LS. 1994. Endocrine effects of prenatal exposure to PCBs, dioxins, and other xenobiotics: implications for policy and future research. Environ Health Perspect 102:676–679.

Blomkvist G, Roos A, Jensen S, Bignert A, Olsson M. 1992. Relationships between PCBs and thyroid hormone levels (e.g., estrogens) (Navas and Segner 1991). J Bone Miner Res 16(10):1812–1820.

Birnbaum LS. 1994. Endocrine effects of prenatal exposure to PCBs, dioxins, and other xenobiotics: implications for policy and future research. Environ Health Perspect 102:676–679.

Birnbaum LS. 1994. Endocrine effects of prenatal exposure to PCBs, dioxins, and other xenobiotics: implications for policy and future research. Environ Health Perspect 102:676–679.

Blomqvist G, Roos A, Jensen S, Bignert A, Olsson M. 1992. Concentrations of sDDT and PCB in seals from Swedish waters. Am. 142:539–545.

Birnbaum LS. 1994. Endocrine effects of prenatal exposure to PCBs, dioxins, and other xenobiotics: implications for policy and future research. Environ Health Perspect 102:676–679.

Blomkvist G, Roos A, Jensen S, Bignert A, Olsson M. 1992. Relationships between PCBs and thyroid hormones and retinol in female and male polar bears. Environ Health Perspect 112:S26–S33.

Brandt M, Siegel MI. 1977. The effects of stress on the cortical bone thickness in rodents. Am J Phys Anthropol 49:31–34.

Colborn T, Vom Saal FA, Soto AM. 1993. Developmental effects of endocrine-disrupting chemicals in wildlife and humans. Environ Health Perspect 101:376–384.

Comiso JC. 2002. A rapidly declining perennial sea ice cover in the Arctic (Letter). Geophys Res Lett 29:1936.

Dartmouth T, Barlow S, Ferguson A, Kavlock RJ, Kaack G. 2002. Global Assessment of the State-of-the-Science of Endocrine Disruptors. Geneva-World Health Organization.
marmoratus in the Norwegian and Russian Arctic. Sci Total Environ 306:159–170.

Lewie, E, Larsen HJS, Larsen S, Johansen GM, Derøcher AE, Lunn NJ, et al. 2004. Does high organochlorine (OC) exposure impair the resistance to infection in polar bears (Ursus maritimus)? Part I: Effect of OCs on the humoral immunity? J Toxicol Environ Health A 67(7):552–562.

Lind PM, Bergman A, Olsson M, Örberg J. 2003. Bone mineral density in male Baltic grey seal. Ambio 32(6):385–388.

Lind PM, Erskine EF, Sahlin L, Eftlind M, Örberg J. 1999. Effects of the antiestrogenic environmental pollutants 3,3′,4,4′,5′-pentachlorobiphenyl (PCB-126) in rat bone and uterus: diverging effects in ovariectomized and intact animals. Toxicol Appl Pharmacol 154:236–244.

Lind PM, Larsson S, Johansson S, Melhus H, Wikström M, Lindh D, et al. 2000b. Bone tissue composition, dimensions and strength in female rats given an increased dietary level of vitamin A or exposed to 3,3′,4,5′-pentachlorobiphenyl (PCB-126) alone or in combination with vitamin C. Toxicology 150:41–51.

Lind PM, Larsson S, Oxlund H, Häkansson H, Nyberg K, Eklund T, Larsson R, et al. 2000a. Change of bone tissue composition and impaired bone strength in rats exposed to 3,3′,4,4′,5′-pentachlorobiphenyl (PCB-126). Toxicology 150:41–51.

Lind PM, Milnes MR, Lundberg R, Bermudez D, Örberg J, Guillette LJ. 2004. Abnormal bone composition in female juvenile American alligators from a pesticide-polluted lake (Lake Apopka, Florida). Environ Health Perspect 112:359–362.

Manolagas SC, Bellido T, Jilka RL. 1995. New insights into the cellular, biochemical and molecular basis of postmenopausal and senile osteoporosis: roles of IL-6 and cellular, biochemical and molecular basis of post-menopausal and senile osteoporosis: roles of IL-6 and osteoclasts. J Bone Miner Res 10(9):1127–1137.

Manolagas SC, Jilka RL. 1995. Organochlorines affect the major androgenic hormone, testosterone, in male polar bears (Ursus maritimus) at Svalbard. J Toxicol Environ Health A 66(21):2119–2139.

Nelson HD. 2003. Postmenopausal osteoporosis and estrogen. Am Fam Physician 68(4):606–615.

Navas JM, Segner H. 1998. Antiestrogenic activity of anthro- genic and natural chemicals. Environ Sci Pollut Res 5:75–82.

Nelson HD. 2003. Postmenopausal osteoporosis and estrogen. Am Fam Physician 68(4):606–615.

Nilsson JA. 1994. Energetic stress and the degree of fluctuating asymmetry—implications for a long lasting, honest signal. Evol Ecol 8(3):248–255.

Norstrom RJ, Belikov S, Born EW, Garner GW, Malone B, Olpienski S, et al. 1998. Chlorinated hydrocarbon contami- nants in polar bears from eastern Russia, North America, Greenland and Svalbard: biomonitoring of Arctic pollution. Arch Environ Contam Toxicol 35(2):354–367.

Oksanen J, Ropstad E, Dahl E, Eie L, Derøcher AE, Wiig O, et al. 2003. Organochlorines affect the major androgenic and estrogenic effects of delta9-tetrahydrocannabinol in the laboratory rat. Toxicol Appl Pharmacol 192:277–284.

Oksanen J, Ropstad E, Dahl E, Derøcher AE, Wiig O, Dahl E, et al. 2004. Organochlorines affect the steroid hormone cortisol in free-ranging polar bears (Ursus maritimus) at Svalbard, Norway. J Toxicol Environ Health A 67:959–977.

Polischuk S, Ramsay M, Norstrom N. 2002. Body burdens and tissue concentrations of organochlorines in polar bears (Ursus maritimus) vary during seasonal fasts. Environ Pollut 118:29–39.

Polischuk SC, Letcher RJ, Norstrom RJ, Ramsay MA. 1995. Preliminary results of fasting on the kinetics of organo- chlorines in polar bears (Ursus maritimus). Sci Total Environ 160/161:465–472.

PMiCrow JHE, Goodman-Gruen D, Sylvania J, Barrett- Connor E. 2002. Retinol intake and bone mineral density in the elderly: the Rancho Bernardo Study. J Bone Miner Res 17(8):1349–1358.

Ramsay MA, Stirling I. 1988. Reproductive biology and ecology of female polar bears (Ursus maritimus). J Zool (Lond) 214:601–634.

Renderer JA, Aulerich RJ, Bursian SJ, Nachreiner RF. 2000a. Prokaryote macrophage and mononuclear periodontal squamous cells in mink fed 3,3′,4,5′-pentachlorobiphenyl (PCB 126). J Vet Diag Invest 12(5):477–479.

Renderer JA, Bursian SJ, Rosenberg DS, Aulerich RJ. 2001. Squamous epithelial proliferation in the jaws of mink fed diets containing 3,3′,4,4′,5-pentachlorobiphenyl (PCB 126) or 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD). Vet Hum Toxicol 43(1):22–26.

Renderer JA, Hocheitin JR, Aulerich RJ, Bursian SJ. 2000b. Proliferation of periodontal squamous epithelium in mink fed 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD). Vet Hum Toxicol 42(5):85–86.

Rosing-Asvid A, Born EW, Kingsley MCS. 2002. Age at sexual maturity of male Baltic grey seal. Ambio 32(6):385–388.

Sandala GM, Sonne-Hansen C, Dietz R, Muir DCG, Valters K, Rothcock DA, Yu Y, Maykut GA. 1999. Thinning of the Arctic sea-ice cover. Geophys Res Lett 26(23):3469–3472.

Sonnenschein C, Dietz R, Leifsson PS, Hjaltul A, Riget FF. 2002. Cadmium toxicity to ringed seals (Phoca hispida)—an epidemiological study of possible cadmium induced nephropathy and osteosteatropy in ringed seals (Phoca hispida) from Qaanaaq in Northwest Greenland. Sci Total Environ 295:167–181.

Söring L, Lunn NJ, Iacoza J. 1999. Long-term trends in the population ecology of polar bears in western Hudson Bay in relation to climatic change. Arctic 52(3):284–296.

Sulc P, Hofbauer LC, Heufelder AE, Roth S, Delmas PD. 2001. Osteoprotegerin serum levels in men: correlation with age, estrogen and testosterone status. J Clin Endocrinol Metab 86(7):3152–3156.

Valentine DW, Soulé M. 1973. Effects of p,p′-DDT on develop- mental stability of pectoral fin rays in the grunion (Leuresthes tenuis). Nat Mar Fish Serv Fish Bull 71:921–925.

Wallgren-Hedlund L, Clasenius AL, Leef P, Thomis M, Philipaerts R, Delvau K, et al. 2002. Association between bone mineral density (DXA), body structure, and body composition in middle-aged men. Am J Hum Biol 14(5):725–742.

Wiig Ø, Derøcher AE, Cronin MM, Skaree JU. 1998. Female pseudohermaphrodite polar bears at Svalbard. J Wildlife Dis 34(4):792–796.

Zakharenko MA, Yakovlev AV. 1990. Skull asymmetry in the Baltic grey seal: effects of environmental pollution. Ambio 19(5):263–269.