1	Electronic Supplementary Information
2	
3	PCBs and PBDEs in environmental samples from King George
4	Island and Ardley Island, Antarctica
5	Pu Wang, Qinghua Zhang*, Thanh Wang, Weihai Chen, Daiwei Ren, Yingming
6	Li and Guibin Jiang
7	State Key Laboratory of Environmental Chemistry and Ecotoxicology, Research
8	Center for Eco-Environmental Sciences, Chinese Academy of Sciences, Beijing
9	100085, China
10	*Corresponding author: E-mail: <u>qhzhang@rcees.ac.cn</u>
11	

12 Materials and methods

13 1. Sampling and shipment

The sampling sites are situated in the Fildes Peninsula at King George Island, west 14 Antarctic, where many international research stations are located including the 15 Chinese Great Wall station, and Ardley Island which is an important settlement for 16 the penguins and migrating birds. During the 26th Chinese Antarctic Expedition in the 17 austral summer between December 2009 and January 2010, twenty-five samples 18 including seven natural soils, three dropping-amended soils, one sediment, six lichens 19 and eight mosses were collected from nine sampling sites that could be reached by our 20 research staff in this area. During the sampling period, the air temperature is around 0 21 ^oC in the Fildes Peninsula and Ardley Island. The soil samples were collected using a 22 stainless steel spade and combined to form a composite sample for the site; the moss 23 and lichen samples were mainly scratched off from rock and ground surfaces; the 24 sediment was obtained using a grab from Great Wall Bay close to the Chinese Great 25 Wall Station. All the samples were stored at -20 °C in the fridge in Chinese Great Wall 26 Station. The sediment was air-dried for several days before the shipment. The samples 27 were sealed in clean plastic bags and packed in a case. After being transported to 28 Chile airport, they were transferred by DHL express to our laboratory in Beijing, 29 China, and then stored at -20 °C. The shipment was finished within 2 weeks. After 30 customs declaration, the package and each sample were checked, and no taint or 31 spoilage was observed for all the samples including the sediment. 32

33 2. Sample extraction and cleanup

For soil and sediment samples, the extraction was performed on Accelerated Solvent Extraction (ASE300, Dionex, USA), and the conditions were as follows: the solvent is DCM: n-hexane (1:1, v:v) (100 mL), the temperature was 150 °C and

S2

pressure was 1500 psi (10.3 MPa) with 7 min for heating and 8 min in the static state. After solvent exchange with DCM and concentrated to 5 mL, the extracts were fractioned using auto-gel permeation chromatography (GPC, AccuPrepTM, J_2 -Scientific; Bio-Beads S-X3 with DCM as the mobile phase) to remove humic substance and sulfur. Then they were loaded to multilayer silica columns packed from the bottom up with 1 g silica gel, 4 g basic silica gel, 1 g silica gel, 8 g acid silica gel, 2 g silica gel and 2 cm anhydrous sodium sulfate, and eluted with 100 mL n-hexane.

For lichen and moss samples, different from the above extraction and cleanup 44 procedures, 3.0 g lichen or moss samples were weighed and the ASE temperature was 45 selected as 100 °C. The extracts were evaporated to dryness for lipid determination, 46 and then dissolved into 50 mL n-hexane. After about 10 g acidic silica was added to 47 remove lipids, the solution was filtered by a column packed with anhydrous sodium 48 sulfate. The filtrate was thereafter concentrated and loaded to a multilayer silica 49 column as previously described, and thereafter through a basic alumina column (6 g 50 basic alumina, 1 cm anhydrous sodium sulfate from the bottom up, eluted with 40 mL 51 *n*-hexane/DCM (1:1, v:v)) and carbon column (1.5 g carbon mixture, 2 cm anhydrous 52 sodium sulfate from the bottom up, eluted with 50 mL *n*-hexane) for further treatment. 53 All the eluates were finally concentrated into 20 µL nonane and then spiked with 54 recovery standard (68A-IS) for the instrumental analysis. 55

- 56 Fig. S1. Relative distribution of PCBs in soil and sediment samples.
- 57 Fig. S2. Relative distribution of PCBs in lichen samples.
- 58 Fig. S3. Relative distribution of PCBs in moss samples.
- 59
- 60
- 61 Table S1. Average concentrations of PCB and PBDE congeners (ranges) in each
- 62 sample type.
- Table S2. Pearson correlation coefficients between the concentrations and TOC in the
- 64 soil and sediment.



Fig. S1. Relative distribution of PCBs in soil and sediment samples.

66

67 Fig. S2. Relative distribution of PCBs in lichen samples.



68

69



70 Fig. S3. Relative distribution of PCBs in moss samples.

72 Table S1. Average concentrations of PCB and PBDE congeners (ranges) in each

73 sample type.74

Compound	Soil (pg g ⁻¹ dw)	Sediment (pg g ⁻¹ dw)	Lichen (pg g ⁻¹ dw)	Moss (pg g ⁻¹ dw)
CB-11	41.7 (19.2-67.9)	5.10	95.4 (45.7-171)	131 (27.5-258)
CB-209	0.60 (n.d.*-2.30)	0.17	3.22 (0.11-16.2)	2.23 (n.d5.68)
Dioxin-like PCBs	,		, , ,	
CB-77	2.31 (0.04-11.1)	0.42	2.19 (0.79-6.25)	1.36 (0.39-1.80)
CB-81	0.12 (n.d0.24)	0.03	0.42 (n.d0.75)	0.56 (n.d0.71)
CB-105	3.84 (0.04-17.4)	1.13	4.02 (0.97-14.2)	3.17 (1.67-5.87)
CB-114	0.27 (0.02-1.05)	0.05	0.60 (0.18-1.30)	0.26 (0.14-0.41)
CB-123	0.30 (0.02-0.90)	0.07	0.30 (n.d0.54)	0.21 (0.12-0.31)
CB-126	0.96 (0.03-2.51)	0.16	1.15 (0.18-2.82)	0.59 (0.26-0.91)
CB-156	2.12 (0.10-8.03)	0.46	2.23 (0.54-4.79)	2.33 (0.93-4.08)
CB-157	0.85 (0.03-2.76)	0.14	0.54 (0.15-1.24)	0.75 (0.27-1.27)
CB-167	1.39 (0.05-4.92)	0.21	0.98 (0.24-2.08)	1.36 (0.34-2.16)
CB-169	1.52 (0.07-3.39)	0.09	0.43 (0.17-0.86)	0.57 (0.31-1.33)
CB-189	0.28 (0.02-1.00)	0.09	0.68 (n.d2.01)	0.53 (0.28-0.87)
Indicator PCBs				
CB-28	25.2 (5.87-78.9)	2.39	50.9 (24.1-76.9)	48.6 (14.3-108)
CB-52	3.67 (0.44-11.5)	1.11	4.05 (1.82-5.72)	3.92 (0.59-6.86)
CB-101	6.64 (0.86-22.5)	2.16	4.82 (3.03-11.2)	5.74 (4.16-8.82)
CB-118	12.0 (0.11-49.1)	2.81	10.9 (2.77-35.5)	9.88 (3.80-19.4)
CB-138	46.5 (n.d174)	3.87	9.29 (1.94-28.1)	17.8 (10.2-31.8)
CB-153	36.3 (n.d143)	4.51	13.4 (3.93-30.3)	27.7 (8.83-59.5)
CB-180	10.2 (0.19-62.5)	1.93	7.26 (1.90-18.2)	20.4 (12.8-40.7)
PBDEs				
BDE-17	1.00 (0.03-5.16)	0.03	0.68 (0.40-0.99)	1.10 (n.d2.11)
BDE-28	3.26 (0.17-15.8)	0.17	1.76 (1.10-2.27)	3.63 (0.87-6.44)
BDE-47	3.84 (1.39-7.37)	1.39	3.93 (2.58-6.48)	4.65 (3.25-6.54)
BDE-66	3.11 (n.d8.04)	n.d.	0.52 (n.d1.12)	1.18 (n.d2.29)
BDE-71	7.25 (n.d17.1)	n.d.	1.05 (0.33-2.01)	1.93 (0.45-5.28)
BDE-85	1.64 (n.d4.07)	n.d.	0.40 (n.d0.64)	1.77 (n.d2.06)
BDE-99	5.84 (n.d13.8)	1.11	2.99 (1.40-4.75)	3.87 (n.d6.26)
BDE-100	0.43 (n.d0.74)	n.d.	0.62 (n.d0.86)	2.05 (n.d2.26)
BDE-138	1.46 (n.d3.79)	n.d.	n.d.	n.d.
BDE-153	1.07 (n.d2.31)	0.48	1.91 (n.d3.77)	2.95 (n.d6.78)
BDE-154	n.d.	n.d.	n.d.	n.d.
BDE-183	n.d.	n.d.	2.18 (n.d3.59)	n.d.
BDE-209	n.d.	n.d.	n.d.	n.d.

75

*n.d., not detected.

Table S2. Pearson correlation coefficients between the concentrations and TOC 76

77 in the soil and sediment.

Commenced	log(POPs) vs. log(TOC)	Commenced	log(POPs) vs. log(TOC)	
Compound	Pearson coefficient	- Compound -	Pearson coefficient	
CB77	0.889**	CB118	0.862**	
CB81	0.779*	CB138	0.766**	
CB105	0.758**	CB153	0.670*	
CB114	0.759**	CB180	0.763**	
CB123	0.891**	CB209	0.727*	
CB126	0.893**	BDE17	0.689*	
CB156	0.917**	BDE28	0.770**	
CB157	0.866**	BDE47	0.697*	
CB167	0.907**	BDE66	0.830**	
CB169	0.564	BDE71	0.844**	
CB189	0.872**	BDE85	0.435	
CB11	0.562	BDE99	0.415	
CB28	0.711*	BDE100	0.894*	
CB52	0.642*	BDE138	-0.082	
CB101	0.695*	BDE153	0.899**	

**. Correlation is significant at the 0.01 level (2-tailed).*. Correlation is significant at the 0.05 level (2-tailed). 79

80