

**Table S1** Analytical techniques and their findings in various research done in various areas

Area	Extraction & Cleanup	Determination	POPs detected	Ref.
Santiago de Compostela (north-western Spain)	<b>Extraction</b> - PLE after the addition of surrogate standards. <b>Clean up</b> - dual layer EZ-POP SPE cartridge	GC-QqQ-MS	PYRs, PAHs, OPPs, OCPs, NDLPCBs, DLPCBs, PBDEs	(19)
Canada	<b>Extraction</b> - 300 mL acetone: hexane (2:1) Homogenization - Polytron. Lipid content - gravimetrically <b>Clean up</b> – diluted (1:1) dichloromethane: cyclohexane ; filtered - 0.45µm polytetrafluorethylene Cleaned up - gel permeation chromatography (S-X3 bio beads;200–400 mesh) <b>Mobile phase</b> - dichloromethane: cyclohexane (1:1, v/v) <b>Flow rate</b> - 5 mL min <sup>-1</sup>	GC coupled to a MicroMass AutoSpec magnetic sector mass spectrometer- Agilent 6890 <b>Column</b> - 30 m DB5 column (0.25 mm x 0.25 µm) coupled (3 m; 0.53 mm retention gap) <b>Temperature</b> - Initial - 80 °C; (2 min); increase 8 °C min <sup>-1</sup> ; 240 °C; final increase - 15 °C min <sup>-1</sup> ; 280°C (5 min). <b>Head pressure</b> - 28 kPa to 173 kPa <b>Carrier gas</b> - Helium. MS- EI positive mode (50 Ev) Source temperature - 250 °C Trap current – 650 µA	Mirex	(20)
Canada	<b>Extraction</b> - Acetone: Hexane (2:1, v/v) <b>Homogenize</b> - Omni Tissue Homogenizer (1min) <b>Centrifuge</b> - Eppendorf centrifuge 5810R (10 min) <b>Clean up</b> - Gel permeation chromatography S-X3 bio beads (200–400 mesh) <b>Mobile phase</b> - 1:1 dichloromethane: hexane (v/v) <b>Flow rate</b> - 5 mL min <sup>-1</sup>	GC coupled to a Waters AutoSpec Premier high-resolution mass spectrometer (Agilent 7890A) <b>Column</b> - 15 m DB-5MS (0.25 mm x 0.10 µm) coupled (5 m, 0.53 mm; deactivated fused silica retention gap) <b>Oven temperature</b> – initially - 80 °C (2 min); increase – 170 °C; 20 °C min <sup>-1</sup> (5.5 min); final increase - 320 °C; 25 °C min <sup>-1</sup> (10 min) <b>Carrier gas</b> - Helium (1.2 mL min <sup>-1</sup> ) <b>MS</b> - EI positive ion mode (36eV) <b>Trap current</b> - 600 µA <b>Source temperature</b> – 250 °C	DDC-CO	(20)

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Lebanon	<b>Extraction</b> - Liquid-liquid extraction; Hexane-Dichloromethane (5:1, v/v) <b>Homogenize</b> – manually (1min) <b>Purify</b> - two steps (1) passing the extract over an Al <sub>2</sub> O <sub>3</sub> glass column with glass frit (22 cm by 20 mm inside diameter) The extract was eluted with Pentane, <b>Evaporated</b> - rotatory Evaporator (2) Fractionation 1.5% (w/w) deactivated silica glass column (15cm by 11 mm inside diameter)	HRGC <b>Detector</b> - <sup>63</sup> Ni electron capture detector <b>Capillary column</b> - DB-608 (30 m by 0.32 mm inside diameter, 0.5 µm film thickness) <b>GC– MS</b> <b>Capillary column</b> SLB-5ms P/N 28471-U (30 m by 0.25 mm inside diameter, 0.25 µm film thickness) <b>Injector temperature</b> – 250 <sup>0</sup> C <b>Purge flow</b> - 2 mL min <sup>-1</sup> (2min) <b>Carrier gas</b> - Helium <b>Flow rate</b> - 1 mL min <sup>-1</sup> <b>Make-up gas</b> - Nitrogen	DDE	(21)
Jinhua, China	<b>QuEChERS extraction</b> modified from (13)	<b>GC-EI/ GC-SIM</b> (Agilent 7890A/5975C-GC/MSD) <b>Column</b> - DB- 17-MS capillary column- (30 m × 0.25 mm × 0.25 µm) <b>Carrier gas</b> - Helium (99.999%) (constant flow rate 1.0 mL min <sup>-1</sup> ) <b>Temperature program</b> - initial (70 °C; 2 min); Increased – (25 °C min <sup>-1</sup> , 150 °C) ; Immediately increased - (200 °C; 3 °C min <sup>-1</sup> ) Finally increased– (8 °C min <sup>-1</sup> ; 290 °C; 6 min.	<i>p,p'</i> -DDE, β-HCH and HCB	(22) (13)

Area	Extraction & Cleanup	Determination	POPs detected	Ref.
Poland	<b>Extraction</b> - Manual; n-hexane/diethyl ether mixture (1:1) <b>Clean up</b> - automated multi-column system PowerPrep (four-channel system-High Capacity Disposable Silica Column, Multilayer Silica Column, Alumina Column, and Carbon Column) <b>Elution solvents:</b> n-hexane, dichloromethane, and toluene (1:1; 2:8; 1:9)	<b>HRGC/HRMS</b> AutoSpec Ultima NT <b>Inlet temperature</b> - 270 °C <b>Column</b> – (60 m × 0.25 mm × 0.25µm) <b>AutoSpec</b> – operated in EI mode, SIR mode <b>Resolution</b> – 10,000 (5%valley) <b>Ion source temperature</b> – 250 °C	PCDD, PCDF, and dl-PCB	(23)
Insular Materno-Infantil University Hospital (Gran Canaria, Canary Islands, Spain; CHUIMI)	<b>Extraction</b> - QuEChERS extraction procedure Acetonitrile saturated in n-hexane <b>Centrifuge</b> - (5,000 rpm, 5 min, 20 °C) <b>Clean up</b> – d-SPE SPE sorbents – PSA	<b>GC-QqQ-MS/MS</b> Fused silica capillary column BPX5 (30 m x 0.25mm x 0.25 µm) <b>Carrier gas</b> - Helium (99.999 %) (flow rate of 1.0 mLmin <sup>-1</sup> ) <b>Oven temperature</b> - Initial - 60 °C (1 min); ramped - 12 °C/min; 210 °C; raised 8 °C/min; 320 °C (6 min) <b>Injector</b> – 270 °C	<i>p,p'</i> -DDE, Hexachloro benzene; <i>β</i> -HCH; <i>γ</i> -HCH; Dieldrin; <i>p,p'</i> -DDD; <i>p,p'</i> -DDT; Dicofol; Mirex; PCBs; PAH (naphthalene, fluorene, phenanthrene, fluoranthene, pyrene, benzo[a]anthracene, chrysene)	(13)

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Yucatan, Mexico	<b>Extraction</b> – SPE Hexane/Toluene (1:1)  <b>Clean up</b> - in LC-Florisil cartridge Conditioned – with 5 mL of n-hexane	<b>GC-ECD</b> – ECD - 63Ni electron capture detector <b>Capillary column</b> - fused silica (30 m × 0.32 mm × 1.0 µm) <b>Carrier gas</b> - Helium (99.999%) (1.5 mL/min with constant pressure) <b>Makeup gas</b> - Ultrahigh purity nitrogen (flow rate - 40 mL/min) <b>Oven temperature</b> - initially at 150 °C (2 min); increased - 210 °C (10 °C min <sup>-1</sup> ); increased - 300 °C (5 °C min <sup>-1</sup> ) <b>ECD temperature</b> - 330 °C; 1.0 nA	OCPs (a- Lindane; b- Lindane; g- Lindane; d- Lindane; Heptachlor; Heptachlor epoxide; Aldrin; Dieldrin; Endrin; Endrin aldehyde; Endosulfan I; Endosulfan II; Endosulfan sulfate; DDE; DDD; DDT	(24)
France	<b>Extraction</b> - Liquid/liquid extraction with pentane. <b>Fat content</b> - gravimetric method <b>Clean up</b> with hexane Isolate OCs – GPC <b>Lipid removal</b> - acid silica; fractionation – florisil; further <b>Purification</b> - celite/carbon columns	<b>GC-HRMS</b> - on electromagnetic sector instruments (JEOL MS 700D or 800D); SIM acquisition mode (PCDD/F, PCB, PBDE, and OC) <b>LC-MS/MS</b> on a triple quadrupole instrument (HBCD isomers)	PCDD/F, PCB, PBDE, and OC HBCD	(25)
Flanders Belgium	<b>SPE</b> on an OASIS HLB (6 mL, 500 mg) cartridge, <b>Purification</b> on an acid silica column	<b>GC-MS</b> <b>Column</b> – HT-8 column (25m x 0.22 mm x 0.25 lm) with MS - EI mode (PCBs, HCB, p,p0-DDE, and p,p-DDT) DB-5 column (15m x 0.25mm x 0.10 lm) MS - ECNI mode (PBDEs and other pesticides)	PCB congeners pesticides HCB, <i>p,p'</i> -DDE, oxychlordan e and <i>b</i> - HCH, dl- PCBs and PCDD/Fs	(26)

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Hong Kong	<b>Extraction</b> - Hot extraction device (Twisselmann extractor) + ethanol/toluene (70/30); 8h; <b>Purify</b> - Butyl methyl <b>Fat extract</b> – dissolved - cyclohexane/ethyl acetate (1:1 v/v) <b>Clean up</b> - gel permeation chromatography (remove fat) <b>Column</b> - (1580 mm x 25 mm x filling level 330 mm) <b>Eluting solvent</b> - Bio-Beads S-X3 + cyclohexane/ethyl acetate (1:1 v/v)	<b>GC/ECD</b> GC (Fisons Mega 2) with two custom-made columns (different polarity) parallel; 1: 30 m PS-088 [97.5% dimethyl-2.5% diphenyl siloxane copolymer], 0.32 mm i.d; 0.32 lm film thickness; 2: 30 m OV-1701-OH, 0.32 mm i.d; 0.25 lm film thickness <b>Confirm</b> - GC–MS (GC: HP 6890/MS: HP 5973; 30 m HP5-MS, 0.25 mm i.d; 0.25 lm film thickness + 2.5 m precolumn <b>Detection mode:</b> MSD-EI GC/MS-EI, SIM mode (PBDE)	HCB, cis-hepta- chlor-epoxide, oxy-chlordane, trans-nonachlor, alpha-HCH, beta-HCH, BDE 47 and BDE 153	(27)

PLE - Pressurized liquid extraction; GC-QqQ-MS/MS - Gas chromatography coupled with a triple quadrupole mass spectrometer; PYRs - Pyrethroids; PAHs - Polycyclic aromatic hydrocarbons; OPPs - Organophosphorus pesticides; OCPs - Organochlorine pesticides; NDLPCBs - Non-dioxin-like polychlorinated biphenyls; DLPCBs - Dioxin-like polychlorinated biphenyls; PBDEs - Polybrominated diphenyl ethers ; EI - Electron impact ionization; DDC-CO - Dodecachlorodimethanodibenzocyclooctane; HRGC - High-resolution gas chromatography; DDE - Dichlorodiphenyldichloroethylene; SIM - Selective ion monitoring; HRMS - High-resolution mass spectrometer; PCDD/Fs - Polychlorinated dibenzo-p-dioxins and dibenzofurans; SPE - Solid phase extraction; ECD - Electron Capture Detector; ECNI - Electron capture negative ionization electron impact; d-SPE - Dispersive solid-phase extraction; PSA - Primary secondary amine; SIR - Selective ion recording; GPC - Gel permeation chromatography; LC-MS/MS - Liquid chromatography coupled to tandem mass spectrometry