

Persistent Organic Pollutants Burden, Trophic Magnification and Risk in a Pelagic Food Web from Coastal NW Mediterranean Sea

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1	Persistent organic pollutants burden, trophic magnification and risk in a pelagic
2	food web from coastal NW Mediterranean Sea
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4	Javier Castro-Jiménez ^{1,2,*} , Daniela Bănaru ¹ , Chia-Ting Chen ¹ , Begoña Jiménez ³ , Juan
5	Muñoz-Arnanz ³ , Geneviève Deviller ⁴ , and Richard Sempéré ¹
6	
7	¹ Aix-Marseille University, University of Toulon, CNRS, IRD, MIO UM 110, Marseille,
8	France.
9	² IFREMER, Laboratory of Biogeochemistry of Organic Contaminants (LBCO), Rue de
10	l'Ile d'Yeu, BP 21105, 44311, Nantes, Cedex 3, France
11	³ Department of Instrumental Analysis and Environmental Chemistry, Institute of
12	Organic Chemistry (IQOG-CSIC), Madrid, Spain
13	⁴ DERAC, 104 Grande Rue, 44240, Sucé-sur-Erdre (Nantes), France
14	
15	*Corresponding author. Phone: +33(0)240374185;
16	E-mail: Javier.Castro.Jimenez@ifremer.fr
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23 Abstract

24 The storage capacity, trophic magnification and risk of sixty-two POPs have been evaluated in a well-characterized pelagic food web (eleven trophic levels) from an 25 impacted area in NW Mediterranean Sea. Our results show the high capacity of the 26 planktonic compartment for the storage of polybrominated diphenyl ethers (PBDEs) and 27 polychlorinated dibenzo-p-dioxins and dibenzofurans (PCDD/Fs), consistent with their 28 estimated low trophic magnification factors (TMF) of 0.2-2.0 (PBDEs) and of 0.3-1.1 29 (PCDD/Fs). Σ PBDEs dominated in the zooplankton size-class 200-1000µm (~330 ng g⁻ 30 ¹ lw, median), whereas Σ PCDD/Fs accumulated preferentially in phytoplankton size-31 class 0.7-200µm (875 pg g⁻¹ lw, median). In contrast, polychlorinated biphenyls (PCBs) 32 were preferentially bioaccumulated in the higher trophic levels (six fish species and two 33 cephalopods) with TMFs=0.8-3.9, reaching median concentrations of 4270 and 3140 ng 34 g^{-1} lw (Σ PCBs) in Atlantic bonito (*Sarda sarda*) and chub mackerel (*Scomber colias*), 35 36 respectively. For these edible species, the estimated weekly intakes of dioxin-like POPs for humans based on national standards overpassed the EU tolerable weekly intake. 37 Moreover, the concentrations of non-dioxin like PCBs in S. sarda were above the EU 38 maxim levels in foodstuffs, pointing to a risk. No risk evidence was found due to 39 consumption of all other edible species studied, also for PBDEs. The integrated burden 40 of POPs in the food web reached ~ 18 μ g g⁻¹ lw, representing a dynamic stock of toxic 41 organic chemicals in the study area. We show that the characterized food web could be a 42 useful and comprehensive 'bioindicator' of the chemical pollution status of the study area, 43 opening new perspectives for the monitoring of toxic chemicals in NW Mediterranean 44 45 coastal waters.

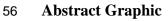
47 Keywords: Environmental exposure, bioaccumulation, flame retardants, dioxins, PCBs,

48 fish, cephalopods, plankton

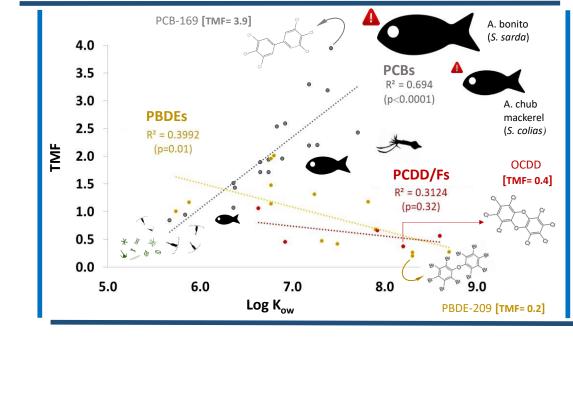
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50 **Synopsis**: The integrated burden of POPs in a pelagic food web (eleven trophic levels) in 51 coastal NW Mediterranean Sea reached ~ $18 \mu g g^{-1}$ lw. A potential risk for humans mostly 52 due to exposure to dioxin-like POPs *via* consumption of Atlantic bonito and chub 53 mackerel was found.

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- 63 1. Introduction
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The chemical contamination of the marine environment is a global and transversal issue with important ecological and socio-economic implications, yet not well quantified nor efficiently managed. ^{1,2} Populations living in coastal areas, as well as marine organisms,

are exposed to multiple sources of chemical contamination, particularly in semi-enclosed 68 69 environments such as the Mediterranean Sea. The Mediterranean is a sensitive environment with the particularity of being a hotspot of both marine biodiversity (up to 70 18% of the world's marine diversity) and chemical contamination (e.g. organic and 71 inorganic contaminants, plastic debris). ³⁻⁵ Among the contaminants impacting this 72 73 marine environment, persistent organic pollutants (POPs) and related chemicals are well documented in non-biotic compartments⁶⁻¹⁰ and different marine organisms.^{5,8,11-17} POPs 74 are organic contaminants of major concern due to their high toxicity, long-range transport 75 and bioaccumulation potential, and their high resistance to degradation.¹⁸ Even if these 76 77 toxic chemicals have been banned by international regulations, such as the Stockholm 78 Convention on POPs (SC), their ubiquity in the marine environment many years after their release may still pose potential risks. 79

Polychlorinated dibenzo-p-dioxins and dibenzofurans (PCDD/Fs), polychlorinated 80 biphenyls (PCBs) and polybrominated diphenyl ethers (PBDEs) are widespread POPs 81 with different environmental sources. While PCDD/Fs can occur as unintentional by-82 products in a number of industrial processes and domestic heating^{19,20}, PCBs are legacy 83 industrial chemicals used in many applications (e.g. lubricants, transformers, oil 84 additives).²¹ Major primary sources for these two families of legacy POPs are considered 85 under control in most parts of the world today.²² However, ongoing primary releases from 86 diffuse sources such as open fires and uncontrolled waste incineration, urban/industrial 87 88 centres, as well as secondary sources such as net re-emissions from environmental reservoirs are still possible.^{21,22} PBDEs were mostly used (and are still in some regions) 89 as flame retardants²³ and, in contrast to PCDD/Fs and PCBs, have been only recently 90 restricted at global scale by the SC.²⁴ These chemicals constitute complex mixtures in the 91 environment to which marine organisms and humans are exposed. Their accurate 92

determination in the biotic compartment represents a very important first step to
apprehend potential risks. However, despite the fact that POPs are often measured in
organisms from different trophic levels (TLs), these organisms do not always belong to
the same food web, making it difficult to properly investigate their trophic magnification,
subsequent risks and more generally the environmental status of marine waters, as
requested by the European Marine Strategy Framework Directive.²⁵

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Here, we report results for PCDD/Fs, PCBs, and PBDEs in 26 biotic samples consisting 100 101 of phyto- and zooplankton, six fish species and two cephalopods, all belonging to the 102 same trophic web from an impacted coastal area, the Bay of Marseille, in the NW Mediterranean Sea. The overall aims of this work were: (1) to characterize a pelagic food 103 104 web representative of the coastal NW Mediterranean, including species consumed by 105 humans, which could help the monitoring and assessment of chemical pollution in the area; (2) to determine the burden and storage capacity of a cocktail of 62 toxic organic 106 107 chemicals in the selected food web; (3) to assess their potential trophic magnification, exposure to humans and risks via consumption of fishery products. 108

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- 110 2. Materials and Methods
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112 *2.1. Study area*

Samples were collected in the Bay of Marseille in the south of France (NW Mediterranean Sea) (Figure S1). This bay is highly impacted by multiple contamination sources, such as the Marseille sewage treatment plant, with its outlet (WWTP Cortiou) located around 10 km south-east of the sampling site,²⁶ and the port, industry, agriculture and tourism, in particular. The Bay of Marseille is naturally influenced by high solar radiation,²⁷

intrusions of the Northern Current on the shelf, ²⁸ and occasionally by intrusions of the
Rhone river plume and the coastal Huveaune River, giving rise to episodic high inputs of
particle and dissolved organic matter. ²⁹⁻³¹ Inputs of a wide diversity of POP families and
related organic contaminants of emerging concern have been reported in the area coming
from the Rhone River, ³²⁻³⁶ the Marseille WWTP at Cortiou and the Huveaunne, ^{35,37-39}
and the atmosphere. ¹⁰ Marine organisms and pelagic food webs in the Bay of Marseille
are certainly exposed to those contaminants (Figure S1).

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126 2.2. *Sampling*

127 Phyto- and zooplankton samples were collected at the Solemio station (43.24°N; 5.29°E, Figure S1) on 15th March 2017 on board the RV ANTEDON II. Phytoplankton and 128 suspended particulate matter (mostly microphytoplankton and detritus particles) were 129 obtained by filtering ~60 L of surface water collected with an inner lining Teflon GoFLo 130 12 L bottle through ten pre-combusted 47mm glass fiber filters (GFFs, 0.7 µm) and 131 treated as one sample. Zooplankton was vertically sampled, from 0 to 50 m depth, using 132 a zooplankton net of 150 µm mesh, and then sequentially separated by using stainless 133 134 steel sieves. Due to the limited amount of sampled material, the various fractions were pooled together into two overall size classes ('Z1': 200-1000µm, and 'Z2': >2000 µm) 135 for POPs analyses. However, all zooplankton size-classes were considered for the food 136 web structure determination (Figure S2). Their groups and species as well as biochemical 137 composition are detailed elsewhere.⁴⁰ Teleosts and cephalopods were sampled by 138 139 professional fishers in the Bay of Marseille near Solemio and specimens were acquired in the Marseille fish market on the same day. Sampled species were: (1) the planktivorous 140 fish bogue (Boops boops, Linnaeus, 1758); (2) the mesopredators Trachurus trachurus 141 (Linnaeus, 1758), Trachurus mediterraneus (Steindachner, 1868), Scomber colias 142

(Gmelin, 1789), the European hake (Merluccius merluccius, Linnaeus, 1758), and the 143 144 cephalopod squid Illex coindetii (Vérany, 1839), Todarodes sagittatus (Lamarck, 1798), 145 and (3) the top predator Atlantic bonito (Sarda sarda, Bloch, 1793). All these fish and cephalopods are commonly caught by local fisheries and consumed in the region. Three 146 147 individuals of different size were considered by species, except for the cephalopod T. sagittatus for which only one specimen was available. In the laboratory, the fresh 148 149 organisms were cleaned, measured and weighed (Table S1). Plankton analyses were made 150 on the entire individuals, while dorsal muscle was sampled for teleosts and mantle for squid. Dissection was done on a clean glass table strictly avoiding contact with plastics 151 152 or any other material which could be a source of organic matter. All the samples were 153 then frozen at -20°C, freeze-dried and ground into a homogenized fine powder prior to 154 analyses (see text S1 for details). We acknowledge that the overall contaminant transfer 155 in the food web will be dictated by the consumption of the entire prey individuals in the 156 ecosystem. However, the choice of the sampling in this study concerned only the muscle for cephalopods and fishes as generally represents the major contaminant exposure route 157 for humans, therefore enabling a risk assessment (section 3.5). Further analyses in higher 158 159 lipid-rich organs such as the liver and gonads may reveal a higher total POP burden and 160 a potentially higher biomagnification.

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162 2.3. Stable isotope and biochemical analyses

163 Three replicates were performed on phytoplankton and zooplankton size classes, while 164 the three individuals of teleosts and cephalopods were considered as replicates for δ^{13} C, 165 δ^{15} N, %C, %N, and total lipid analyses. Stable isotope measurements were performed 166 with a continuous-flow isotope-ratio mass spectrometer coupled to an elemental analyzer 167 in LIENSs Laboratory, La Rochelle University (France). Details for total lipid determinations are presented in Text S1. For fish samples, δ^{13} C values were corrected for the effects of lipids when the C/N >3.5.⁴¹ TLs were estimated using the zooplankton 200-300 µm size class (composed mostly of small herbivorous copepods) as 'baseline' with trophic position defined as 2.^{40,42} A classical enrichment factor of 3.4‰ between prey and predator was applied except for the top predator *S. sarda* for which a value of 2.4‰ was used ⁴³ (See Text S1 for all details).

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175 2.4. POPs analysis

176 All samples were freeze-dried and zooplankton and fish samples homogenized by using 177 a ball mill mounting stainless steel cells. The corresponding amounts of sample (0.14-0.25 g dw, Table S1) were spiked with ¹³C-labeled standards of PCDD/Fs, PCBs, and 178 PBDEs prior to Soxhlet extraction (24h) with a n-hexane: DCM (9:1) mixture. Extracts 179 were rota-evaporated and cleaned-up using the automated sample preparation system 180 DEXTech+.¹⁴ The six indicator PCBs (_{ID}PCBs), the eight mono-*ortho* dioxin-like PCBs 181 182 (DLPCBs) and PBDEs (27 congeners) were collected in a first fraction, while the 183 seventeen 2,3,7,8-substituted PCDD/Fs along with the four non-ortho _{DL}PCBs were obtained in a second fraction, summing up a total of 62 POPs. Final extracts were rota-184 185 evaporated to ~ 1 mL, transferred to vials, and dried under a gentle nitrogen steam. Fractions were reconstituted in a few microliters of the respective PCDD/F, PCB, and 186 PBDE ¹³C-labeled injection standards prior to instrumental analysis. Quantification was 187 carried out by isotopic dilution according to 1613 US EPA method ⁴⁴ on a gas 188 chromatograph (Trace GC ultra, Thermo Fisher Scientific) coupled to a high-resolution 189 190 mass spectrometer (DFS, Thermo Fisher Scientific). Details can be found in Text S1.

191 2.5. *Quality assurance / quality control (QA/QC)*

Field blanks for phytoplankton samples (n=2), consisting of baked GFFs which were 192 193 placed in the freeze-dryer and then processed with the samples, were performed. 194 Analytical laboratory blanks (n=6) accounting for all the sample processing steps were also performed. After sampling, GFFs were wrapped in aluminium foil again and stored 195 196 in the dark at -20°C. Laboratory blank levels were generally low for the three POP classes compared to their concentrations in the samples, averaging from not detected (n.d) to 25 197 pg (PCBs), from n.d. to 0.13 pg (PCDD/Fs), and from n.d. to 7 pg (PBDEs) depending 198 on the congener and sample (Table S3). Laboratory blanks showed lower or similar levels 199 200 to field blanks, so no contamination occurred during sampling, storage and analysis. 201 Results were blank-corrected by subtracting the average field and laboratory blanks 202 values for phytoplankton and the rest of samples, respectively. Chromatographic peaks were only considered when the ratio between the two monitored ions was within $\pm 15\%$ 203 204 of the theoretical value, and the signal-to-noise (S/N) ratio>3 (limit of detection, LOD). 205 Limits of quantification (LOQs) corresponded to S/N≥10. Calibration curves were daily checked. Median LODs (n=26) ranged from 0.2 to 2.3 pg (PCBs), from 0.1 to 0.8 pg 206 (PCDD/Fs) and from 0.3 to 14.0 pg (PBDEs) depending on the compound and sample 207 208 (Table S4). Median method recoveries varied from 60 to 111 % (PCBs), from 79 to 102 209 % (PCDD/Fs) and from 61 to 102 % (PBDEs) (Table S5). Results were corrected by 210 recoveries.

211 2.6. Data analysis

Non-parametric tests (Kruskal–Wallis and Wilcoxon rank-sum) were used in order to investigate significant differences among pollutant levels. The effect of multiple comparisons was considered by applying the false discovery rate method. Non-detected values were imputed to the LOD/2 to run these tests (Text S2 and Table S6). The software employed was STATA/SE 16.1. However, the concentration sums presented in sections

3.3.1 and 3.3.2 consider only the detected values, and the concentrations expressed as 217 218 World Health Organization-toxic equivalents (WHO₀₅-TEQ) presented in section 3.5 for 219 _{DL}POPs are reported as upper bound values (i.e. the n.d values were substituted by the corresponding LODs before the blank correction) to facilitate their comparison with 220 221 available threshold values. The sample BB-1 was excluded from the data set for PBDEs due to suspected specific contamination. Trophic magnification factors (TMFs) for 222 individual POPs were calculated using the slope (m) of the linear regression between the 223 log₁₀ of the lipid-normalized contaminant concentration in the food web organisms and 224 their corresponding TLs (calculated as indicated above) ^{45, 46} (section 3.4.). For PBDEs 225 and PCDD/Fs, which exhibited more n.d values, only congeners with >30% detection 226 frequency and detected in at least 7 different TLs were considered for the TMF 227 calculation. In order to derive the estimated weekly intake (EWI) of $_{DL}$ POPs and \sum PBDEs 228 229 (section 3.5.), the weekly consumption data for France for demersal (*M. merluccius*) and pelagic (rest of species) fish and cephalopod (222, 132 and 9.2 g/week, respectively)⁴⁷ 230 231 were multiplied by the WHO₀₅-TEQ (PCDD/Fs + $_{DL}PCBs$) and the $\sum PBDEs$ (only detected values) concentration, respectively, measured in the corresponding species and 232 divided by 70 Kg (average human bodyweight, bw), as previously described. ¹⁵ Risk 233 quotients (R), were calculated as the concentration in the sample, and its corresponding 234 235 EWI, to threshold values, legally binding or not, derived by international authorities to protect humans consuming fishery products and aquatic ecosystems. 236

237

238 **3.** Results and discussion

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240 3.1. Food web characterization

The stable isotope ratios of the species and size classes analysed in the Bay of Marseille pelagic food web are presented in Figure S2A and Table S7. Both δ^{13} C and δ^{15} N are

significantly different between species and/or size classes (Kruskal-Wallis, p < 0.0001). 243 244 Zooplankton > 2000 μ m mainly constituted of gelatinous organisms has the lowest stable 245 isotope ratios related to their filter feeding regime based on small pico (0.2-2 μ m) and nano (2-20 µm) phytoplankton. These small size classes of phytoplankton are supposed 246 247 to be the main source of the zooplankton food web in the North-Western Mediterranean area ⁴⁸ but they could not be sampled for this study in sufficient quantities for POPs 248 249 analyses. Particles issued from the Cortiou WWTP with low stable isotope ratios are also 250 susceptible to be consumed by the different zooplankton size classes and decrease their stable isotope ratios during particulate rain or hydrodynamic conditions. ^{26,42} Micro-251 phytoplankton size class may represent the prey of many planktivorous fish such as B. 252 253 boops, and all the juveniles of the studied fishes and squid. Herbivorous zooplankton 200-300 µm had also low stable isotope value and were considered as baseline for the 254 estimation of the TL of the other consumer groups. The $\delta^{15}N$ ratios increased between 255 low TL groups (phyto- and zooplankton) and high TL groups (fish and squid, TL > 3) 256 consistent with their TL position (Figure S2B/Table S7). Predators such as the fish S. 257 sarda and squid T. sagittatus showed the highest stables isotope ratios related to their 258 259 high TL (5.3 and respectively 4.2).

260

261 3.2. Distribution of POPs and storage capacity in the food web

Overall, PCBs exhibited a high detection frequency (DF) of 73-100% for most congeners (Figure S3), and dominated total POP concentrations from zooplankton to top predators, accounting for 69% to around 100% of the \sum POPs (Figure 1A). PBDEs showed a generally lower DF with most congeners in the range 0-60%, although a certain number of congeners such as BDEs-28, -47, -49, -71, -99, -100 and -154 were highly detected (DF= 84-100%). PBDEs dominated only in phytoplankton, reaching 64% of total POPs

concentrations, while accounting for 0.3 - 31% of the total contaminant stock in the rest 268 269 of the samples. PCDD/Fs levels were below LODs in many cases and DF was mostly in 270 the range 0-40% and only 2,3,7,8-TCDF (88 %) and 1,2,3,4,6,7,8-HpCDD (96%) were highly present. The contribution of PCDD/Fs to the total POPs burden in the studied food 271 web was minor, representing < 0.5% of the total POPs. These contaminants accumulated 272 preferentially in the lowest trophic levels (LTL) like PBDEs (Figure 1B). LTL, 273 274 considered as the planktonic compartment (i.e. phytoplankton 0.7-200 µm, zooplankton 200-1000 μ m and zooplankton > 2000 μ m), seemed to play a predominant role driving 275 276 the food web burden and storage capacity of Σ PBDEs and Σ PCDD/Fs exhibiting 12-fold and 5-fold higher median concentration than in higher trophic levels (HTL), respectively 277 (Figure 1C). This trend was followed by individual PBDEs and PCDD/Fs too (Figure S4-278 S5) showing similar or significantly higher (p < 0.01 - 0.04, Kruskal-Wallis) median 279 concentrations in LTL. In contrast, the \sum PCB burden in the food web may be dominated 280 281 mostly by HTL showing a significantly 9-fold higher median concentration (p=0.02, Kruskal-Wallis) than the LTL (Figure 1B). Individual PCB congeners generally followed 282 the same trend except for CBs-28 and -52, which exhibited similar or significantly higher 283 284 median concentrations (p=0.03, Kruskal-Wallis) in LTL as PBDEs and PCDD/Fs (Figure S6). In other words, the lipid-normalized cumulative concentrations of PBDEs and 285 PCDD/Fs in the phytoplankton and the two zooplankton size classes were similar or 286 significantly higher than the cumulative concentrations of the 21 fish and 2 cephalopod 287 individuals studied, pointing to the high capacity of the planktonic compartment for the 288 accumulation and storage of these two POPs families. In contrast, fishes and cephalopods 289 290 represent efficient biotic storage compartments for PCBs.

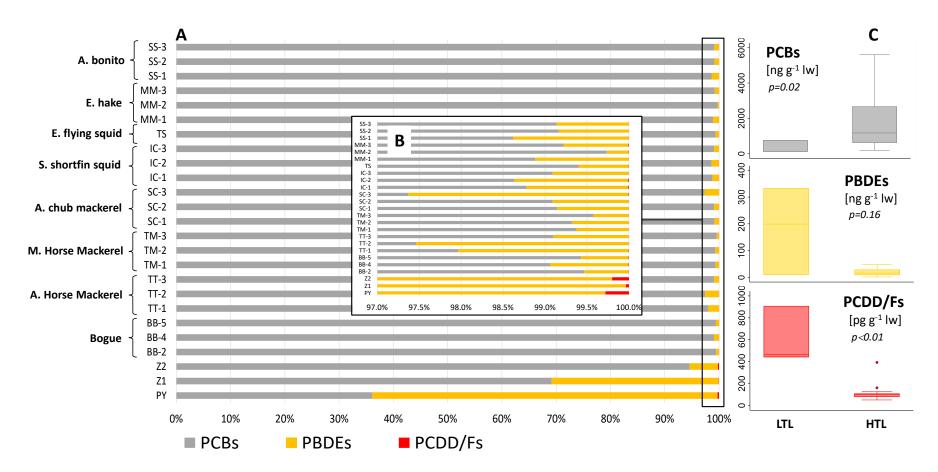


Figure 1. POPs relative burden in the food web (A) with a zoom showing the higher % of PCDD/Fs found in the planktonic compartment (B), and box plots (C) of the lipid normalized total concentrations of Σ PCBs, Σ PBDEs (ng g⁻¹ lw) and Σ PCDD/Fs (pg g⁻¹ lw) in the lowest trophic levels (LTL) considered as the planktonic compartment for the purpose of this comparison (i.e. *PY=phytoplankton 0.7-200 µm, Z1=zooplankton 200-1000 µm and Z2=zooplankton > 2000 µm, n=3*) and the higher trophic levels (HTL) considered as the other organisms from the food web (fishes and cephalopods, n=22) Kruskal-Wallis test p-value. *BB=B. boops; TT=T. trachurus; TM=T. mediterraneus; SC=S. colias; IC=I. coindetii; TS=T. sagittatus; MM=M. merluccius; SS=S. sarda.*

301 *3.3.1. Overall concentrations*

 Σ_{18} PCB concentrations varied from 3.3 to 496 ng g⁻¹ dw (from 113 to ~5600 ng g⁻¹ lipid 302 weight - lw) in the different individuals from the studied food web. The total PCB load 303 was dominated by \sum_{ID} PCBs, accounting for 83-92% of \sum_{18} PCBs with concentrations 304 ranging from 3.0 to 454 ng g⁻¹ dw (~100 to 5130 ng g⁻¹ lw). \sum_{DL} PCBs represented only 305 between 8 and 17% of the total PCBs with levels between 0.3 and 42 ng g^{-1} dw (13 to 470 306 ng g⁻¹ lw). \sum_{26} PBDE concentrations varied from 0.08 to 25 ng g⁻¹ dw (1.0 to 333 ng g⁻¹ 307 308 lw) whereas three orders of magnitude lower \sum_{17} PCDD/F concentrations were measured, ranging from 0.3 to 110 pg g⁻¹ d.w (3.4 to 875 pg g⁻¹ lw). Results for individual PCBs, 309 PBDEs and PCDD/Fs in each sample are presented in Tables S8-10. In addition, in order 310 311 to facilitate data comparability, POPs concentrations are also expressed on a wet weight (ww) basis (Table S11). 312

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314 *3.3.2. Levels and patterns in the different organisms*

315 Although concentrations of a number of POPs, mostly PCBs and PBDEs, were found significantly different among the different TLs in a first data analysis (p < 0.0 - 0.04, 316 Kruskal-Wallis) (Table S6), when comparing by pairs of TLs (Wilcoxon rank-sum test) 317 and correcting by the effect of multiple comparisons (section 2.6, Text S2), these 318 319 differences were not statistically significant. These analyses are certainly hampered by 320 the limited amount of data. However, some trends are worth discussing. Although some bioaccumulation was observed in the 200-1000 µm zooplankton for the sum of PCBs, the 321 species exhibiting the highest median \sum_{18} PCB concentration was the Atlantic bonito (S. 322 sarda) (4270 ng g⁻¹ lw), presenting the highest position in the food web (i.e TL=5.3), 323

324	followed by the Atlantic chub mackerel (S. colias) (3140 ng g^{-1} lw). The European flying
325	squid (T. sagittatus) and the Mediterranean horse mackerel (T. mediterraneus) were also
326	among the highest concentrations with 2670 ng g ⁻¹ lw (only one sample) and 2280 ng g ⁻¹
327	lw (Table 1). Similar trends were found for individual PCBs, except for CB-28 and -52
328	which accumulated preferentially in the planktonic compartments with no clear trends in
329	the other TLs (Figure S7). A consistent PCB pattern was observed in the whole food web,
330	with PCB-153 as the most abundant congener in all TLs (27-42% of the median
331	\sum_{18} PCBs), except for the TL 1 (phytoplankton) where PCB-101 and -52 were more
332	abundant. The predominance of PCBs-101 and -52 has been reported in the dissolved
333	water phase in other NW Mediterranean coastal areas.49 Likely their preferential
334	bioconcentration at the very first TLs could explain this differential pattern found in
335	phytoplankton. PCB-118 was the most abundant $_{DL}PCBs$ contributing ~53 – 74% of the
336	median \sum_{DL} PCBs (Figure S8).

Table 1. Median concentrations (only detected values) of indicator, dioxin-like, and total PCBs (\sum_{ID} PCBs, 339 \sum_{DL} PCBs, \sum_{18} PCBs, respectively), total PBDEs (\sum_{26} PBDEs) and total PCDD/Fs (\sum_{17} PCDD/Fs) in the 340 different trophic levels of the food web under study. Values are expressed in dry, wet and lipid weights 341 (dw, ww, lw, respectively). Concentrations are in ng g⁻¹ except for PCDD/Fs (pg g⁻¹). 342

Median concentrations*	$\sum_{ID} PCB$ [ng g ⁻¹]		$\sum_{DL} PCB$ [ng g ⁻¹]			$\sum_{18} PCB$ $[ng g^{-1}]$			\sum_{26} PBDEs [ng g ⁻¹]			$\sum_{17} \text{PCDD/Fs}$ $[\text{pg g}^{-1}]$			
Wedian concentrations.															
Foodweb Species	dw	ww	lw	dw	ww	lw	dw	ww	lw	dw	ww	lw	dw	ww	lw
Phytoplankton (0.7-200 µm)	12.5	2.5	99.2	1.7	0.3	13.8	14.2	2.8	113.0	25.0	5.0	199.0	109.9	22.0	874.6
Zooplankton (200-1000 µm)	39.3	7.9	653.0	5.7	1.1	95.4	45.1	9.0	748.3	20.1	4.0	333.3	23.7	4.7	393.4
Zooplankton (>2000 µm)	3.0	0.6	112.7	0.3	0.1	12.9	3.3	0.7	125.6	0.2	0.04	6.9	7.0	1.4	265.9
Boops boops	38.4	9.2	629.4	6.2	1.5	102.2	44.7	10.7	732.3	0.5	0.1	5.3	2.1	0.5	28.8
Trachurus trachurus	55.7	11.1	858.5	5.6	1.1	79.2	61.3	12.3	937.6	1.0	0.2	19.5	0.7	0.1	9.2
Trachurus mediterraneus	180.2	36.0	2023.4	18.9	3.8	256.0	200.3	40.1	2279.4	0.8	0.2	15.6	0.9	0.2	16.2
Scomber colias	203.0	58.9	2734.2	19.1	5.5	286.4	222.1	64.4	3140.8	1.9	0.6	28.7	2.6	0.8	38.8
Merluccius merluccius	36.8	8.8	538.1	3.9	0.9	45.9	41.1	9.9	584.0	0.3	0.08	4.8	1.6	0.4	23.8
Illex coindetii	69.3	16.6	1229.4	8.1	2.0	155.5	77.5	18.6	1384.9	0.8	0.2	14.9	6.2	1.5	152.2
Todarodes sagittatus	181.8	43.6	2355.8	24.4	5.9	316.3	206.2	49.5	2672.1	1.2	0.3	16.1	6.5	1.6	84.5
Sarda sarda	444.0	106.5	3903.9	41.6	10.0	367.5	485.7	116.6	4271.4	4.2	1.0	37.1	3.9	0.9	43.3

343

"only one sample was available for phytoplankton, the two zooplankton size clasess and the cephalopod *Todaroa*

345 Median \sum_{26} PBDE concentrations clearly dominated in the 200-1000µm zooplankton (~330 ng g⁻¹ lw) and phytoplankton (199 ng g⁻¹ lw), whereas \sum_{17} PCDD/Fs accumulated 346 preferentially in phytoplankton (875 pg g⁻¹ lw) followed by 200-1000 µm zooplankton 347 (~390 pg g^{-1} lw). The \sum PBDE predominance in zoo- and phytoplankton was driven by 348 the higher contribution of BDE-209 (70-75% of the detected Σ PBDEs). The high 349 \sum_{17} PCDD/F levels found in the LTL were driven by the higher abundance of 350 octachlorodibenzo-p-dioxin (OCDD) accounting for the 64-80% of the detected 351 \sum PCDD/Fs (Figure S9). Indeed, the predominant of OCDD/Fs in zooplankton has been 352 reported in other marine food webs.⁵⁰ Both BDE-209 and OCDD are highly hydrophobic 353 POPs with log octanol-water partition coefficients (log Kow) of 8.7 and 8.2, respectively, 354 indicating high partition potential on hydrophobic components such as lipids or organic 355 carbon-rich particles. The combination of the following factors may explain this 356 357 observation: The 200-1000 µm plankton size classes in the study area, mostly composed of copepods, have been reported to have a high concentration of protein and lipid.⁴⁰ 358 359 Moreover, the composition of smallest size class (0.7-200 µm) is dominated by 360 microphytoplankton and detritus (mostly from the Marseille WWTP effluents and Rhone inputs), and small copepod stages^{26,40,42}, proving active surfaces for the accumulation of 361 highly hydrophobic contaminants. A previous study showed important atmospheric 362 inputs of particle-bound PBDEs (~550 ng m⁻² y⁻¹) and PCDD/Fs (~15 ng m⁻² y⁻¹) in the 363 area, also with a clear predominance of BDE-209 and OCDD in the atmospheric particle 364 patterns.¹⁰ This atmospheric loading may constitute a relatively constant and direct source 365 of these congeners to surface waters, facilitating their way to the LTL of the food web. 366 367 Regarding HTLs, S. sarda and S. colias presented the highest Σ PBDE median

and, *T. mediterraneus* exhibiting median concentrations in the range 16-20 ng g^{-1} lw

368

concentrations of 37 and 29 ng g⁻¹ lw, respectively, followed by *T. trachurus*, *T. sagittatus*

(Table 1). Although individual PBDEs generally followed this trend too (Figure S10), 370 371 some differences were found in the PBDE patterns. In the group of species showing higher Σ PBDE median concentrations (i.e. S. sarda, S. colias, T. trachurus, T. 372 373 mediterraneus, and T. sagittatus), the PBDE pattern was characterized mostly by the predominance of BDE-47 and -100 (32-46% and 13-31% of the detected Σ PBDEs, 374 respectively) and a lower contribution of PBDE-209 (4-16% of the detected Σ PBDEs) 375 (Figure S11A). The predominance of these congeners, particularly BDE-47, has been 376 reported for other marine organisms.^{14,17,51} Furthermore, the patterns in *M. merluccius* 377 378 and I. coindetii, two out of the three species showing the lowest median values, showed a more even relative distribution of BDE-47 and BDE-209, accounting for the 24-32 and 379 380 32-39% of the detected Σ PBDEs, respectively (Figure S11B). These are piscivorous and 381 invertivorous species feeding mainly on pelagic species. Interestingly, the pattern found in B. boops showed a similar BDE-209 contribution as M. merluccius and I. coindetii but 382 lower relative contribution of BDE-47 and higher % of BDE-99, resembling instead that 383 of phytoplankton and 200-1000µm zooplankton (Figure S11B,C). The fact that B. boops 384 is essentially zooplanktivorous⁴⁰ could partially explain this pattern. The 200-1000µm 385 386 zooplankton have been reported as the zooplankton class-size with the highest energy content in the study area, making it preferable prey for zooplanktivorous fish such as the 387 bogue.⁴⁰ However, a differential metabolic capacity of *B. boops* could also contribute to 388 389 the observed pattern. Species-specific differences in metabolic rates of PBDE in fish has been already reported.⁵² A higher median BDE-99/BDE-100 ratio (reported as an 390 indicator of fish capacity to metabolize PBDEs)^{49, 53} for *B. boops* (0.9) was observed in 391 comparison to the combined ratio for *M. merluccius* and *I. coindetii* (0.2), suggesting a 392 less efficient capacity of *B. boops* to metabolize PBDEs. For comparison, the Σ PBDE 393 concentrations in HTL organisms in this study (0.02-1.0 ng g⁻¹ww, Table S11B) were 394

similar or slightly lower to those reported in fish from other Mediterranean locations. Values ranging from 0.07 to 3.4 ng g⁻¹ww were measured in wild fish from the Adriatic, Ionian and Tyrrhenian Seas (E Mediterranean)¹⁷ and average concentrations ranging from ~1.0 to 3.9 ng g⁻¹ ww were reported for fish and seafood collected in Valencian markets (Spain, NW Mediterranean)⁵⁴. The fact that the authors of these studies reported the upper bound concentrations and that other fish species were also studied could explain an important part of the concentration variability.

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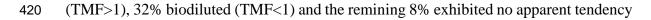
Regarding HTL, Σ PCDD/Fs accumulated preferentially in the cephalopods exhibiting 403 median concentrations of 152 and 84 pg g^{-1} lw in the Southern shortfin squid (*I. coindetii*) 404 and European flying squid (T. sagittatus), respectively. The pattern in these organisms 405 was characterized by a predominance of OCDD (around 50% of the detected \sum PCDD/Fs) 406 407 and was quite similar to that of the planktonic compartment (Figure S12A). Individual congeners followed the same concentrations trend, except for 2,3,7,8-TDCF which also 408 accumulated in S. colias, B. boop, S. sarda and T. trachurus (Figures S13), as reflected 409 410 too in their respective patters (Figure S12B) with this congener accounting from ~60 to 411 76% (median) of the detected \sum PCDD/Fs. *M. merluccius* and *T. mediterraneus* showed a 412 comparable contribution of 1,2,3,4,6,7,8-HpCDD, 2,3,7,8-TDCF, and OCDFs 413 representing 30-40% of the detected \sum PCDD/Fs depending on the congener and species 414 (Figure S12C).

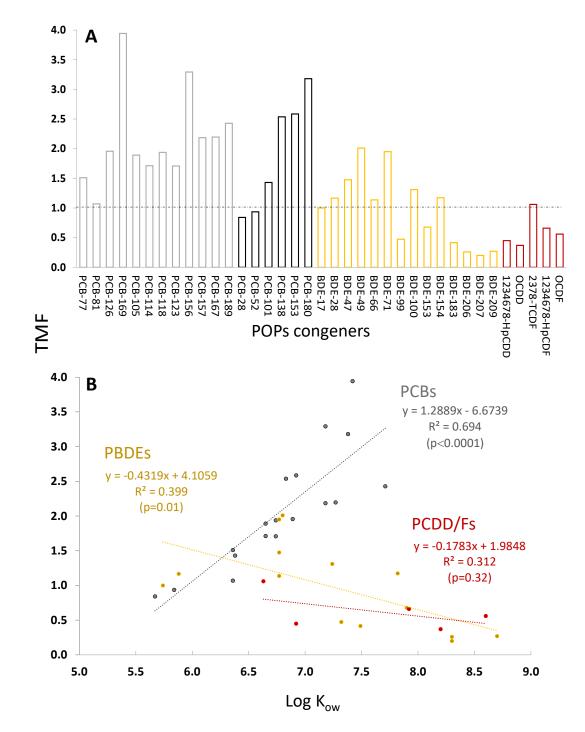
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416 3.4.Trophic magnification

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TMFs were calculated for the eighteen PCBs, fourteen PBDEs and five PCDD/Fs (Figure
S14, Table S12). The majority of POPs (60%) with calculated TMFs bioamplified





421 $(TMF~1)^{45}$ (Figure 2A).

422

423 Figure 2. Trophic magnification factors (TMFs) for individual PCBs, PBDEs and PCDD/Fs (A) and their 424 relationship with the POP's log K_{ow} (B) for the NW Mediterranean Sea food web. Only congeners with DF 425 > 30% and expanding at least along 7 different TLs were considered. The horizontal dotted line indicates 426 TMF=1. Light and dark grey, yellow and red indicates DLPCBs, IDPCBs, PBDEs and PCDD/Fs, 427 respectively. The standard errors and confidence intervals of the slopes used to derive the TMFs can be 428 consulted in Table S12.

PCBs showed the highest biomagnification potential (>80% congeners with TMF>1) with 430 431 TMFs ranging from 1.4 (PCB-101) to 3.9 (PCB-169). Our results indicate that even if 432 PCB-153, one of the most studied PCBs in aquatic organisms, exhibited a lipidnormalized concentration of 4-5 orders of magnitude higher than PCB-169 and PCB-126 433 434 (Table S8), its estimated trophic magnification potential ($TMF_{CB-153} = 2.6$) is around 35% 435 weaker than that for PCB-169 and only 24% higher than that of PCB-126 (TMF=1.9). 436 PCBs-126 and -169 are indeed the most toxic _{DL}PCBs (toxic equivalent factor, TEFs=0.1 and 0.03, respectively).⁵⁵ This may have implications for the appropriate environmental 437 438 exposure assessment to the most toxic PCBs in the study area, which may be overlooked by only studying the most abundant PCBs, which is often the case, and warrants further 439 research. Only PCB-28 (TMF=0.8) and -52 (TMF=0.9) did not biomagnify, consistent 440 with their higher abundance in the food web LTL (section 3.3.2). A significant positive 441 correlation ($R^2=0.69$, p<0.0001) was found between the TMF_{PCBs} and the log K_{ow} (Figure 442 443 2B) confirming the general higher trophic magnification for the more hydrophobic PCBs in this Mediterranean food web, as previously reported for other aquatic ecosystems⁴⁶. 444 This biomagnification potential is consistent too with the higher PCB concentrations 445 generally found in the Atlantic bonito (Tables 1, S7 and Figure S7). 446

447

In contrast to PCBs, only 50% of PBDE congeners biomagnified, exhibiting generally lower TMFs ranging from 1.1 (BDE-66) to 2.0 (BDE-49). The other congeners showed a trophic dilution in the food web with TMFs varying from 0.2-0.7 (Figure 2A, Table S12). A negative significant correlation (R^2 =0.40, p=0.01) was found for TMF_{PBDEs} and the log K_{ow} indicating a lack of biomagnification of the heaviest PBDEs like BDE-206, 207 and 209 among others (Figure 2B), consistent with their higher predominance in the planktonic compartment. It is worth mentioning that no or little trophic magnification

(0.5 < TMF < 1.5) was found for most of the PBDEs considered in the European Quality 455 Standard for PBDEs (i.e. BDE-28, -47, -99, -100, -153, -154) in fishes (EOS_{biota})⁵⁶. 456 Moreover, a generic biomagnification factor (BMF) of 5 is used for those PBDEs to cover 457 their biomagnification from lower trophic levels to fish species ⁵⁷, which is much higher 458 459 than our filed-derived TMFs. PCDD/Fs showed TMFs ranging from 0.4 to 1.1 indicating mostly trophic dilution, with a non-significant negative correlation ($R^2=0.31$, p=0.32) 460 461 between TMF_{PCDD/Fs} and log K_{ow}. This lack of significant correlation with log K_{ow} has also been reported in other aquatic food webs. ⁵⁸ Similarly to PBDEs, a BMF=10 is 462 considered for PCDD/Fs and _{DL}-PCBs to account for their biomagnification in aquatic 463 food webs ⁵⁹ which is much higher than the TMFs measured in this study. These fact 464 465 raises the importance of using site and food web specific biomagnification values in order to perform a more reliable site-specific risk assessment instead of applying generic default 466 467 values.

468

Overall, our results indicate that the POPs biomagnification in the studied food web is 469 strongly congener and species dependent. The fact that POPs congeners with similar Kow 470 471 showed distinct bioaccumulation patterns, particularly the more hydrophobic PCBs, 472 PBDEs and PCD/Fs indicated that lipophilicity was not the only factor explaining their trophic magnification. This differential behaviour has also been reported in other aquatic 473 food webs.⁵⁸ Indeed, the interplay of the specific contaminant bioavailability and 474 metabolic transformation may have driven their final biomagnification. The observed 475 trophic dilution of higher hydrophobic PBDEs in wild fish could be mostly attributed to 476 the metabolic debromination of HMW PBDEs as previously reported. ⁵² However, the 477 biodilution of higher hydrophobic PCDD/Fs, exhibiting negligible metabolization rates 478

- in most cases, may be better explained by a reduced membrane permeability due to their
 larger molecular size compared to the lower chlorinated congeners. ⁵⁰
- 481

Field-derived TMFs for the studied POPs in marine food webs are scarce in the literature 482 483 for comparison. In addition, TMFs in marine systems have been reported to be highly variable and a number of factors such as type, structure and length of the food web and 484 485 the site study specificities and season of the year, may have an important influence on the final TMF value ^{60,61}. Therefore, the comparability of TMFs must be carefully interpreted. 486 TMFs of 1.3 and 1.2 were reported for PCB-153 and BDE-47, respectively, in a study 487 conducted in the Gulf of Lion during the period 2004-2006 and in a shorter foodweb⁶⁰. 488 489 These values are slightly lower than those calculated for the same contaminants in this study (TMF_{PCB-153} = 2.6 and TMF_{BDE-47} = 1.5). The fact that we studied a longest food 490 491 web, including a higher top predator (A. bonito), could explain our higher values. TMFs 492 within the same range were reported for a pelagic food web in the western Scheldt estuary (Netherlands)⁶² for BDE-28, -47, and -100. This work highlighted too the trophic dilution 493 of BDE-209 with a very similar TMF of 0.2. However, they found BDE-99 494 495 biomagnifying (TMF=2.4) in their studied food web, while a TMF=0.5 was found for the 496 same congener in our study. The reported predominance of BDE-99 in the western Scheldt estuary organisms ⁶², likely linked to historical local sources, could explain this 497 difference, since this congener was not predominant in our food web. TMFs for PCBs 498 499 and PBDEs within the same range were reported for a pelagic food web in coastal Atlantic Ocean⁶³. However, our TMFs for these two families of POPs were generally lower than 500 501 the average (± standard deviation) TMFs reported for a global compilation also including fresh water ecosystems, which ranged from 1.7 ± 1.1 to 8.1 ± 6.9 for PCBs and from 0.3 502 ± 0.3 to 3.2 ± 1.9 for PBDEs ⁶¹, although a large variability was reported too (Figure S15). 503

To the best of our knowledge, we report here the first field-derived TMFs for PCDD/Fs in a Mediterranean food web, which are very similar to those reported in other aquatic ecosystems⁶¹ (Figure S15).

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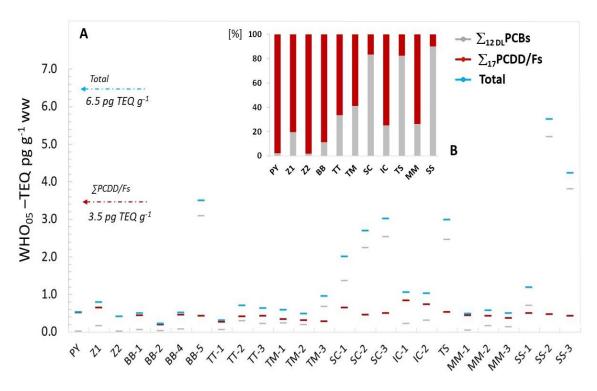
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508 3.5. Toxicity potential and risk assessment

510 *3.5.1.* _{DL}POPs

511 Concentrations expressed as upper bound WHO₀₅-TEQ were derived for _{DL}PCBs and the 512 2,3,7,8-PCDD/Fs in the food web organisms using the latest available TEFs⁵⁵ in order to 513 study their toxicity potential against humans. Concentrations ranged 0.008 - 5.2 and 0.2 514 - 0.8 WHO₀₅ TEQ pg g⁻¹ ww for Σ_{DL} PCBs and Σ_{17} PCDD/Fs, respectively, resulting in 515 TEQ_{Total} (Σ_{29} dioxin-like POPs) of 0.2-5.7 pg g⁻¹ ww (Figure 3A, Table S13).

516



517

Figure 3. Calculated WHO₀₅-TEQ (pg g⁻¹ ww) in all samples from the pelagic food web in coastal NW Mediterranean Sea (A) and median relative contribution (%) of \sum_{12DL} PCBs (grey) and \sum_{17} PCDD/Fs (red) to the total WHO₀₅-TEQ (PCBs+PCDD/Fs) in blue (B), except for PY, Z1, Z2 and TS, which are single values. The horizontal dotted lines show the EU concentration thresholds in sea foodstuffs and wild fish (see text for details). *PY=phytoplankton 0.7-200 µm; Z1=zooplankton 200-1000 µm; Z2=zooplankton >* 2000 BB=B. boops; TT=T. trachurus; TM=T. mediterraneus; SC=S. colias; IC=I. coindetii; TS=T. sagittatus; MM=M. merluccius; SS=S. sarda.

The species S. sarda, T. sagittatus and S. colias exhibited the highest median TEQ_{Total} 526 values of 4.2, 3.0, and 2.7 pg g⁻¹ ww, respectively (Table S13). These TEQs were driven 527 by the predominant contribution of _{DL}PCBs (82-90%) in those species (Figure 3B). 528 Median TEQ_{Total} values in the remaining trophic levels varied from 0.4 to 1.1 pg g^{-1} ww. 529 530 PCDD/Fs exhibited a generally higher contribution (53-98%) on these species, except for some isolated samples such as BB-5 and TM-3, which show a higher concentration of 531 PCB-126, increasing the relative contribution of \sum_{DL} PCBs TEQ on those samples. The 532 533 \sum_{DL} PCB TEQ concentrations measured in the studied food web were within the range or higher than those reported for other fish species in the Mediterranean. For example, levels 534 535 of 0.4-1.2 (Sardina pilchardus), 0.08-0.4 (Engraulis encrasicolus) and 0.07-0.27 (B. *boops*) TEQ pg g⁻¹ ww have been recently reported.¹⁴ Our concentrations in fish samples, 536 excluding those of the top predator Atlantic bonito, ranged from 0.02 to 3.1 TEQ pg g⁻¹ 537 ww for \sum_{DL} PCB, with values varying from 0.05 to 3.1 TEQ pg g⁻¹ ww for *B. boops*. Our 538 539 \sum_{DL} PCB TEQ values were also higher than those reported for chub mackerel (*Scomber* japonicas) (0.002 -0.008 TEQ pg g⁻¹ ww) and horse mackerel (T. trachurus) (0.004 -540 0.016 TEQ pg g⁻¹ ww) caught in the Adriatic Sea.⁶⁴ For comparison, our \sum_{DL} PCB levels 541 in T. trachurus varied from 0.05 to 0.3 TEQ pg g⁻¹ ww (Table S13). A recent study 542 measured dioxin-like POPs in Mediterranean bluefin tuna (Thunnus thynnus), reporting 543 0.7 and 1.9 TEQ pg g⁻¹ ww for \sum_{DL} PCB and \sum PCDD/Fs, respectively.⁶⁵ Our results in 544 the Atlantic bonito ranged from 0.7 to 5.2 and from 0.4 - 0.5 TEQ pg g⁻¹ ww for \sum_{DL} PCB 545 and \sum PCDD/Fs, respectively. However, the concentrations reported for the bluefin tuna 546 were lower bound values (non-detected values were considered as zero). 547 These comparisons are only indicative due to the associated variability related to the different 548 549 species considered, sampling year and season and the quantification strategies adopted in each specific study. 550

At European level, two pieces of legislation, Commission Regulation 1259/2011⁶⁶ and 551 Directive 2013/39/EU⁵⁶, set a maximum level for _{DL}POPs in seafood and marine 552 organisms aiming at protecting human health via consumption of fishery products (but 553 also aquatic top predators from secondary poisoning for the EQS_{biota}). A threshold level 554 of 6.5 pg TEQ g⁻¹ ww for the \sum_{17} PCDD/Fs + \sum_{12DL} PCBs in fish muscle and fishery 555 products, and wild fish (EQS_{Biota}) is established. In addition, regulation 1259/2011 also 556 set a threshold value of 3.5 pg TEQ g⁻¹ ww for \sum_{17} PCDD/Fs only. The concentrations 557 measured in the various species considered did not exceed any of these thresholds, 558 although certain S. sarda and S. colias individuals (i.e. SS-2-3 and SC-3) approached the 559 6.5 pg TEQ g⁻¹ ww threshold for total _{DL}POPs (Figure 3A) with risk quotients (R) between 560 0.5 and 0.9 (R>1 indicating risk) (Table S14). The _{DL}POPs EWIs considering fish and 561 cephalopod consumption data in French population (section 2.6) were calculated for each 562 sample and varied from 0.1 to 11 pg TEQ Kg⁻¹ bw week⁻¹. All individuals of S. colias and 563 S. sarda exceeded the tolerable weekly intake (TWI) of 2 pg TEQ Kg⁻¹ bw week⁻¹ set by 564 the European Food Safety Authority⁶⁷ with R values ranging from 1.1-5.3 and 1.9-2.8 for 565 S. sarda and S. colias, respectively (Table S14). The major contributors to the EWI and 566 subsequent risk for humans associated with the consumption of S. colias and S. sarda 567 were _{DL}PCBs, particularly PCB-126. 568

Moreover, Directive 1259/2011 sets a maximum level (75 ng g⁻¹ ww) for the sum of the six non-dioxin-like PCBs in seafood, also referred as _{ID}PCB in this work. The \sum_{ID} PCB concentrations in the studied food web were for most samples below that threshold (Table S11), varying from 0.6 to 59 ng g⁻¹ ww. However, some *S. colias* individuals again approached the threshold (R=0.8 for SC-1-2) and two out three samples of the top predator *S. sarda* (i.e. SS-2-3) exceeded it with concentrations between 106-109 ng g⁻¹ ww (R=1.4, Table S14) pointing to a risk for humans to non-dioxin-like PCBs due to 576 consumption of this last species. It has been reported that non-dioxin-like PCBs, which 577 are weak aryl hydrocarbon receptor agonists, can modulate the overall toxic potency of 578 2,3,7,8-TCDD and related compounds⁵⁵ and therefore their co-occurrence cannot exclude 579 cumulative effects including synergic harmful effects.

- 580
- 581 *3.5.2. PBDEs*

Regarding PBDEs, the EQS_{Biota} of 0.0085 ng g⁻¹ ww (as Σ_6 PBDEs including BDE-28,-582 47,-99,-100,-153,-154) based on human toxicity⁵⁶ was used for a first risk evaluation. 583 This threshold value was exceeded in every single individual in all TLs (Table S14), 584 585 indicating a risk for humans consuming the edible species, again pointing to S. sarda (R=54-97) and S. colias (R=15-65) as the most 'risky' fish. However, some authors argue 586 that the existing EQS_{Biota} for PBDEs is extremely low leading to environmental 587 concentrations exceeding the threshold value in most study cases.^{51,68} Moreover, for 588 589 OSPAR assessment, there are still questions on the reliability and appropriateness of some of the thresholds set as EQS_{biota}, based on human protection, to support 590 environmental assessment. It has been recently proposed to adopt the Canadian Federal 591 Environmental Quality Guidelines (FEQGs) based only on wildlife protection.⁶⁹ 592 593 Comparing our concentrations with the proposed FEQGs, which have been set for a number of individual PBDE congeners (i.e BDE-28, 47, 99, 100, 153, 154, 183 and 209), 594 there is no indication of apparent risk (Table S15). However, it should be noted that the 595 596 FEQGs are derived for whole organisms, and PBDE were measured in muscle in this study. The Σ PBDEs EWI in France ranged from 0.02 to 1.9 ng Kg⁻¹ bw week⁻¹, the highest 597 value corresponding to S. sarda, followed by S. colias (0.7-1.2 ng Kg⁻¹ bw week⁻¹). For 598 comparison, an average estimated intake of 0.14 ng Kg⁻¹ bw d⁻¹ via fish and seafood 599 consumption for adults has been reported for Valencia (Spain) in the NW Mediterranean 600

Sea⁵⁴, corresponding well with our higher end estimate (0.17 ng Kg⁻¹ bw d⁻¹). There is no 601 602 TWI for Σ PBDE to the best of our knowledge, but the US Agency for Toxic Substances and Disease Registry and the US Environmental Protection Agency have established 603 acute/sub-chronic Minimal Risk Levels⁷⁰ and chronic oral reference doses⁷¹, respectively, 604 for some of the most abundant PBDE congeners (i.e BDE-47, BDE-99, BDE-153, BDE-605 209). These health standards ranged from 3 to 7000 ng Kg⁻¹ bw d⁻¹ depending on the 606 607 congener, showing that there is no risk associated with these congeners for humans 608 consuming these Mediterranean fishery products.

609

610 Our study demonstrates that the edible species Atlantic bonito (S. sarda) and Atlantic 611 chub mackerel (S. colias) bioaccumulated the highest concentrations of POPs which may 612 pose a risk for humans consuming these species caught and /or marketed in the Gulf of 613 Lion (NW Mediterranean Sea) mostly due to _{DL}POPs exposure. The cephalopods 614 southern shortfin squid (I. coindetti) and European flying squid (T. sagittatus) showed a high potential for the accumulation of PCDD/Fs, however no indications of risk can be 615 observed according to our evaluation. An in-depth review of available thresholds for 616 617 PBDEs concentration in biota and the determination of tolerable daily intakes for more 618 congeners are urgently needed to provide a more comprehensive risk assessment. We 619 consider the calculated EWI as a lower-end estimate of their total exposure, since their 620 intake through other food items is not considered and external exposure routes were not 621 accounted for in this evaluation. For example, the reported median outdoor inhalation intake in the study area for _{DL}POPs (0.028 pg TEQ Kg⁻¹ bw week⁻¹) and Σ PBDEs (0.015 622 ng Kg⁻¹ bw week⁻¹) could range up between 0.4 - 1.3% and 0.8-2.1% their respective total 623 intakes. These results warrant further research on the overall intake rates of the three POP 624 625 families by humans and their cumulative risks in this impacted region of NW

Mediterranean Sea. Our results would be highly valuable to set up-to-date and comparable 626 concentration thresholds in biota considering trophic level normalization as 627 recommended by EU guidance⁷² and OSPAR.⁷³ Besides, the characterized food web, as 628 a whole, may be a useful and site-specific comprehensive 'bioindicator' of the chemical 629 pollution status, representing a dynamic stock of toxic organic chemicals in the study 630 area. A lower-end estimation of its integrated POPs burden in the study season, calculated 631 considering the sum of the median concentrations in all TLs, reached ~ $18 \mu g g^{-1}$ lw. This 632 value could be considered a benchmark for further studies using the same food web, 633 opening new perspectives for the monitoring of toxic chemicals in Mediterranean coastal 634 635 waters.

636

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639

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651

652 Supporting information available

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Additional data and details on the sampling and analytical procedures, QA/QC,
concentration of individual congeners, TMFs among others are available free of charge
via the Internet at http://pubs.acs.org/

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