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1 **Spatial variability and sources of platinum in a contaminated harbor – tracing coastal**
2 **urban inputs**

3 Melina Abdou^{1,2*}, Jörg Schäfer¹, Teba Gil-Díaz¹, Mary-Lou Tercier-Waeber², Charlotte
4 Catrouillet¹, Francesco Massa³, Michela Castellano³, Emanuele Magi³, Paolo Povero³, Gérard
5 Blanc¹

6 ¹*University of Bordeaux, UMR CNRS 5805 EPOC, 33615 Pessac, France*

7 ²*University of Geneva, Dept. of Inorganic and Analytical Chemistry, 1211 Geneva 4,*
8 *Switzerland*

9 ³*University of Genoa, DISTAV-DCCI, 16132 Genoa, Italy*

10 **Corresponding author: melina.abdou@unige.ch*

11

12 **Environmental context**

13 Emerging contaminants such as platinum (Pt) are continuously released into aquatic
14 environments. However, due to analytical challenges related with their accurate quantification,
15 little information exists on their geochemical cycles in coastal areas. This paper examines the
16 potential use of Pt as a tracer of anthropogenic inputs in a model industrial/urban site (Genoa
17 Harbor, Italy) and provides a first estimate on dissolved Pt baseline concentrations in the
18 northern Mediterranean coast.

19

20 **Abstract**

21 Biogeochemical cycles including processes controlling platinum (Pt) distribution remain
22 widely unknown in aquatic environments, especially in coastal systems. Dissolved Pt
23 concentrations in coastal seawater (Pt_D) and in Suspended Particulate Matter (SPM, Pt_P) were
24 measured, together with master variables comprising dissolved oxygen, dissolved and
25 particulate organic carbon, chlorophyll-a, turbidity, and ammonium levels, along two
26 longitudinal profiles in the industrial Genoa Harbor (northwest Italy). Concentrations and
27 spatial distribution of Pt_D and Pt_P levels reflect distinct concentration gradients that were
28 attributed to different Pt sources such as hospital, domestic and industrial wastewater,
29 atmospheric deposition, and/or road run-off. Concentrations reaching up to $0.18 \text{ ng.L}^{-1} Pt_D$ and
30 $14 \text{ ng.g}^{-1} Pt_P$ reflect the impact of Pt urban inputs in coastal sites. These first data highlight
31 considerable anthropogenic contamination in the confined harbor compared to the proposed
32 reference value for the western Mediterranean surface seawater measured at external sites.
33 Identified correlations between Pt levels and human pollution signals suggest the potential use
34 of Pt as a new tracer of anthropogenic inputs that can be applied in other urbanized coastal
35 systems. Biogeochemical processes inducing changes in partitioning and fate of Pt in coastal
36 seawater reflect spatial variability, highlighting the need for comprehensive environmental
37 monitoring at appropriate spatial scale.

38 **Keywords:** Platinum; Technology Critical Element; emerging contaminant; seawater;
39 suspended particulate matter; distribution coefficient; urban tracer; coastal environment

40

41 **1. Introduction**

42 Platinum (Pt) is a Technology Critical Element (TCE) and an emerging contaminant of growing
43 concern according to evidences of anthropogenic disturbance on its global biogeochemical
44 cycle (Rudnick and Gao, 2003; Cobelo-García et al., 2015). Among its specific properties, Pt
45 is an excellent catalyst used in industrial processes, such as petroleum chemistry, but more
46 importantly in vehicle catalytic converters to reduce gaseous pollutants in exhaust fumes.
47 Strongly increasing Pt demand for this technology results in growing anthropogenic Pt inputs
48 into the environment, leading to a global increase of Pt concentrations in all environmental
49 compartments including soils, atmosphere and aquatic systems (e.g. Rauch et al., 2006; Rauch
50 and Morrison, 1999; Schäfer et al., 1999). However, other anthropogenic Pt sources (potentially
51 implying the release of different Pt species to the environment) may occur in aquatic
52 environments including Pt-derived anti-cancer drugs (e.g. Vyas et al., 2014).

53 Coastal areas are facing high anthropogenic pressure due to the increasing density of human
54 population in such attractive zones (Small and Nicholls, 2003). Monitoring of trace metal
55 concentrations in these sites is therefore essential to detect potential contamination and prevent
56 associated adverse consequences on the ecosystem, especially for emerging contaminants.
57 Platinum concentration in seawater is difficult to quantify precisely, in particular in less
58 contaminated sites, because of extremely low environmental levels (sub ng.L^{-1} ; Mashio et al.,
59 2017). Existing studies on marine coastal areas, including saline estuaries, report dissolved Pt
60 concentrations ranging from 0.05 to 0.30 ng.L^{-1} (Cobelo-García et al., 2013, 2014a; Goldberg
61 et al., 1986; Mashio et al., 2016, 2017; Obata et al., 2006; Turetta et al., 2003; van den Berg
62 and Jacinto, 1988). A small number of studies report Pt concentrations in Suspended Particulate
63 Matter (SPM) ranging from ~ 2 to 10 ng.g^{-1} in the Lérez Estuary and from ~ 0.2 to 1 ng.g^{-1} in
64 the Gironde Estuary (Cobelo-García et al., 2013, 2014a). Variations in Pt concentrations and

65 possible Pt enrichment in those coastal sites have been attributed to Pt inputs from human
66 activities, especially in systems receiving anthropogenically-impacted freshwater inputs, as observed in the Tokyo Bay (e.g. Obata et al., 2006). Several studies
67 have addressed bioaccumulation of Pt in wild marine bivalves, such as oysters and mussels
68 (Abdou et al., 2016; Neira et al., 2015). However, there is a lack of data on the comprehensive
69 characterization of the factors controlling the behavior of Pt in coastal seawater, including its
70 partitioning between suspended particles and seawater, and huge uncertainties remain
71 concerning its transport, bioavailability and fate in the coastal ocean (Cobelo-García et al.,
72 2013).

74 The Mediterranean Sea is a semi-enclosed system under ever-growing anthropogenic pressure
75 along its northwestern coast, comprising several highly populated cities with extensively
76 urbanized and industrialized areas (Martín et al., 2009). Considering the possible changes in
77 seawater chemistry due to additional anthropogenic inputs, previous studies have addressed
78 environmental quality in historical and industrial harbors such as the Toulon Bay or the Genoa
79 Harbor, reflecting intense pollution by metals, (e.g. Oursel et al., 2013; Tessier et al., 2011),
80 metal-organic components (Pouget et al., 2014) and nutrients (Ruggieri et al., 2011). The
81 Genoa Harbor on the northwest coast of Italy (Ligurian Sea) is one of the most important ports
82 of the Mediterranean Sea, known for heavy contamination due to a variety of human activities.
83 For many years, industrial effluents were the major cause of water pollution in this site, before
84 the closure of the main heavy industries adjacent to the harbor during the 1990s. Nowadays,
85 discharges from sewage treatment plants and from creeks account for generally high levels of
86 nutrients, organic matter and faecal pollution in the harbor waters (Ruggieri et al., 2011).

87 The present work provides first information on Pt levels and spatial distribution in this major
88 Mediterranean harbor combined with physical-chemical parameters. Such findings may help to

89 improve our understanding on Pt sources, other than the commonly assumed influence from
90 catalytic converter emissions, and Pt cycling in an anthropogenically impacted coastal site. The
91 objectives are therefore: (i) to quantify Pt levels and spatial distribution in a marine system
92 under strong anthropogenic influence, involving different Pt sources, (ii) to characterize the
93 factors controlling Pt distribution and partitioning between seawater and suspended particles
94 considered a prerequisite for a better understanding of its biogeochemical cycle in coastal
95 environments, and (iii) to investigate the potential use of Pt as an innovative tracer of human
96 inputs (traffic, wastewaters, industrial activities) in coastal environments.

97

98 **2. Experimental**

99 2.1. Study area and sampling strategy

100 *The coastline of Genoa*

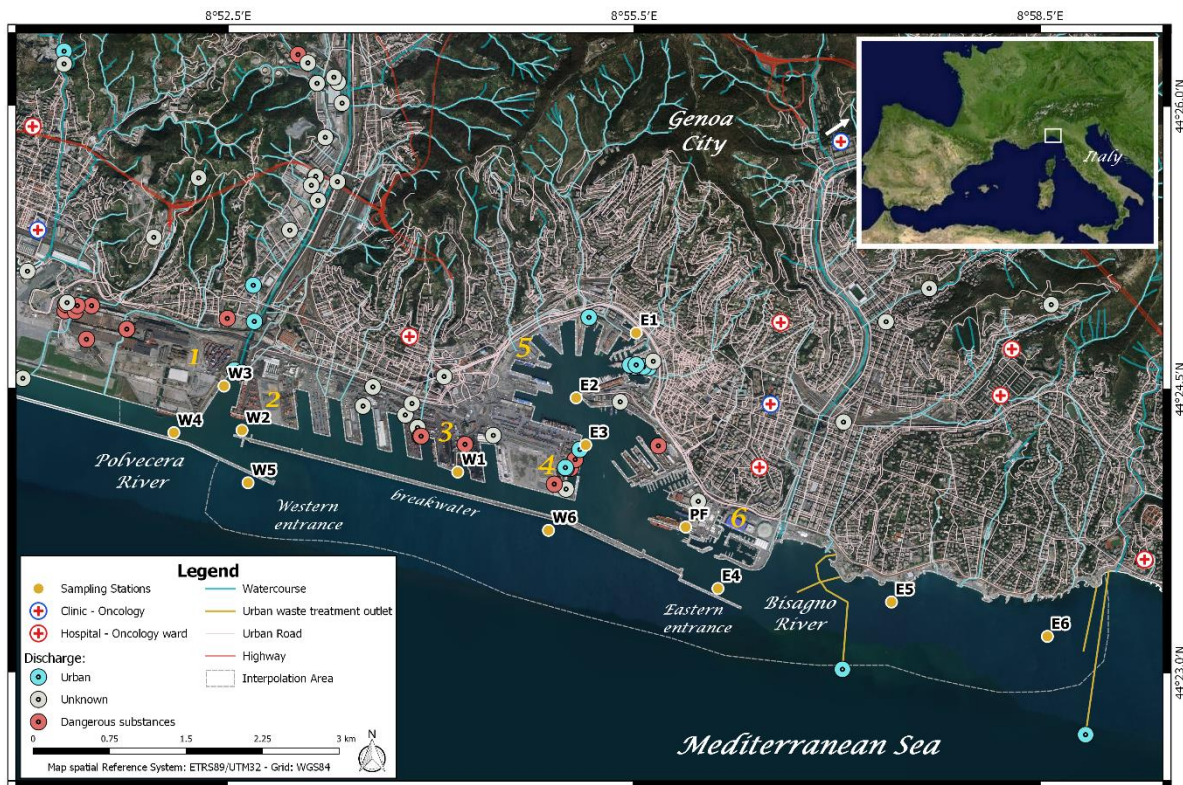
101 The Genoa Harbor is an artificial harbor on the northwest coast of Italy (44°24'39.8"N
102 8°55'44.8"E). The port infrastructures extend over approximately 7 million m² along about
103 20 km of coastline, protected by sea defenses and artificial breakwaters. The innermost, natural
104 part of the Genoa Harbor is the so-called Old Port hosting a ferry terminal, several marinas and
105 receiving urban treated sewage effluents (Ruggieri et al., 2011; Fig. 1). Outside the Old Port a
106 wide, less confined area protected by artificial breakwaters extends to the east and to the west,
107 hosting diverse industrial activities and receiving discharges from a coal-fired power plant,
108 treated urban wastewater and untreated surface run-off (Ruggieri et al., 2011). In addition,
109 important dredging has affected the area in the last decade (Cutroneo et al., 2012). Two
110 seasonally active rivers discharge freshwater and treated wastewater into the harbor area: the
111 Polcevera River to the west and the Bisagno River to the east (Fig. 1; catchment surface area of
112 93 km² and 140 km², respectively; Cutroneo et al., 2017)). Both rivers, characterized by

113 irregular and harsh streamflow fluctuations, run through the city of Genoa and are surrounded
114 by dense urban structures hosting a variety of commercial and industrial activities.

115 *Sampling campaigns*

116 Two longitudinal profiles were performed to collect surface seawater and SPM samples inside
117 and outside the Genoa Harbor on-board the Research Vessel MASO (3rd – 6th April 2017;
118 Fig. 1). The first profile was carried out from the inner part of the port towards the eastern
119 entrance (sampling sites E1 to E6; Fig. 1). This represents a supposed decreasing contamination
120 gradient covering the strongly confined Old Port (sites E1 and E2), a less confined zone inside
121 the breakwaters (site E3), the eastern Inlet (site E4) close to the mouth of the Bisagno River and
122 two sites outside the Harbor, directly in contact with open Mediterranean water, i.e. the Ligurian
123 Current (sites E5 and E6). This profile is performed along the less industrialized section of the
124 harbor. The second profile addressed the western part of the Genoa Harbor (sampling sites W1
125 to W6, Fig. 1) hosting coal deposits, a terminal for oil products (no storage tanks in the harbor),
126 and receiving power plant discharges and inputs from the Polcevera River transporting sewage
127 treatment plant effluents (Ruggieri et al., 2011).

128 Eastward profile was performed on April 3rd under north-northeast wind with an average speed
129 of 16 km.h⁻¹ (Table 1; timeanddate.com). Westward profile was done the following day with
130 weaker wind (~ 9 km.h⁻¹) originating from south-southeast. Due to complex harbor shape,
131 station bottom depth differs according to the location. Within the Old Port, site E1 has a bottom
132 depth of 6 m (Table 1). Other sites within the breakwaters, show average depth of 11 m while
133 it is of 16 m for E4 at the eastern inlet. We observe average depth of 14 m for external sites E5
134 and E6, and 26 m for W5 and W6.



135
 136 **Fig. 1:** Sampling sites in the Genoa Harbor. Two sampling profiles were performed with an
 137 eastward profile (sites E1 to E6) and a westward profile (sites W1 to W6). Sampling site PF
 138 (PlatForm) corresponds to the CNR Platform. Numbers correspond to main urban/industrial
 139 activities with 1: Steel mill, 2: Bulk containers, 3: Coal power plant, 4: Oil storage, 5: Ferry
 140 terminal, 6: Shipyards/marinas. Red crosses indicate hospitals with oncology ward with the
 141 Oncology Clinics highlighted in blue. Colored circles show discharges from various sources.
 142 Geospatial data provided by the Ligurian Administrative Region via own Geoportal - WFS
 143 service (<https://geoportal.regiosne.liguria.it/>). Satellite image provided by ESRI©
 144 (ArgGIS/Imagery)

145
 146 Additional samples were taken at the CNR (National Research Council) Observation Platform
 147 (site PF, Fig. 1) on April 6th 2017 at three different times of the day (10 am, 12 pm, 2 pm). The
 148 bottom depth of this sampling site is of 6 m (Table 1). This sampling site is located at the limit
 149 of the confined area of the harbor, supporting diverse harbor activities such as navigation, ship
 150 construction and wrecking industries. Main hospitals with oncology ward of Genoa are located
 151 rather close to this area, including an Oncology Clinic (~ 3 km).

152 *Sample collection*

153 Seawater and SPM samples were collected at all sites (i.e. on-board the R/V MASO or from
154 the CNR Platform) by pumping surface water (~ 5 m depth) with a purpose-built low voltage
155 (12 V) peristaltic pump connected to Teflon tubing. The latter was rinsed with seawater,
156 accounting for more than three times the tubing volume (seawater pumping time of 10 min)
157 prior to sample collection. Seawater was collected into acid-washed 500 mL Teflon bottles
158 (Nalgene®), previously rinsed with pumped water at each site. Water samples for analyses of
159 dissolved Pt (Pt_D) were filtered immediately through 0.2 μm polycarbonate filters
160 (Nucleopore®) with a filter-syringe (Sartorius®) into acid-washed 60 mL Teflon bottles
161 (Nalgene®), previously rinsed with an aliquot of the filtrate, then acidified to pH = 1 (36.5–
162 38 % HCl Baker Instra) and stored in the dark at 4 °C pending analysis. Water aliquots from
163 the 500 mL Teflon bottle were collected to quantify ammonium (NH_4^+) and dissolved organic
164 carbon (DOC) concentrations. The samples for NH_4^+ analysis were filtered as aforementioned
165 into pre-cleaned 15 mL polypropylene (PP) tubes (Kartell®), and stored frozen pending
166 analysis. Aliquots for DOC were filtered through pre-combusted (500 °C, 4 h) glass-fiber filters
167 (0.7 μm , Whatman® GF/F) using a pre-cleaned (RBS detergent, thoroughly rinsed with ultra-
168 pure deionised water, Milli-Q® water) and pre-combusted glass syringe. Filtrates were
169 collected in pre-cleaned and pre-combusted glass bottles, acidified (HCl 0.3 % v/v; Baker,
170 Analyzed) and then stored in the dark at 4 °C until analysis.

171 Particulate samples were collected for analyses of particulate platinum (Pt_P) and particulate
172 organic carbon (POC). In both cases, the sampling bottle was carefully shaken and seawater
173 was placed in a pre-rinsed filtration unit covered by a lid. For Pt_P , SPM were collected on pre-
174 weighed Teflon filters (47 mm, 0.45 μm FHLC filter, Merck Millipore®), whereas for POC,
175 SPM were collected on pre-combusted and pre-weighed glass-fiber filters (0.7 μm , Whatman®
176 GF/F). Once in the laboratory, both filters were then oven-dried (48 h at 50 °C) to constant

177 weight, re-weighed (to determine SPM content) and kept in the dark at room temperature
178 pending analysis. A field blank was performed for all sampled parameters and processed in the
179 same way as seawater samples but using Milli-Q water.

180

181 2.2. Analytical methods

182 *Platinum in seawater*

183 Seawater samples were analyzed by Adsorptive Cathodic Stripping Voltammetry (AdCSV) as
184 described in Cobelo-García et al. (2014b). Measurements were carried out using a μ Autolab
185 Type III potentiostat (Metrohm®) connected to a polarographic stand (Metrohm® 663 VA
186 Stand) equipped with three electrodes: (i) a hanging mercury drop electrode (HMDE, the
187 working electrode), (ii) a silver/silver-chloride (Ag/AgCl) reference electrode, and (iii) a glassy
188 carbon auxiliary electrode. A polytetrafluoroethylene (PTFE) voltammetric cell served in all
189 experiments and the potentiostat was controlled using the NOVA 2.1® software. Elimination
190 of organic matter by UV oxidation was performed by placing sample aliquots in capped Teflon
191 bottles with overnight irradiation using two 64 W UV lamps (NIQ 60/35 XL, Heraeus) under a
192 fume hood. Aliquots (10 mL) of UV-digested sample were pipetted into the voltammetric cell
193 together with 30 μ L of 3.3 mM formaldehyde (37–41 % Analytical Reagent Grade, Fisher
194 Chemical®), 30 μ L of 0.45 mM hydrazine sulfate (Analytical Reagent Grade, Fisher
195 Chemical®), and 300 μ L of sulfuric acid (H₂SO₄, 93–98 % Trace metal grade, Fisher
196 Chemical®). Platinum concentrations were quantified by standard internal addition method,
197 adding a controlled amount of a Pt stock solution freshly prepared daily from a mono-
198 elementary Pt standard solution (1000 μ g.mL⁻¹ PLASMACAL, SCP Science®). We applied a
199 deposition time of 300 s and experimental parameters as described elsewhere (Cobelo-García
200 et al., 2014b). In the absence of Certified Reference Material (CRM) for dissolved Pt in

201 seawater, analytical quality of the voltammetric procedure was evaluated by means of Pt spiked
202 CRM seawater quantification (CASS-6, NRCC) giving recoveries > 80% and precision
203 expressed as Relative Standard Deviation (RSD) lower than 10 % (n = 5) at the 0.5 ng.L⁻¹ range.
204 The detection limit (D.L.) for dissolved Pt measured by AdCSV (calculated as 3 x blank
205 standard deviation, n = 20) was estimated to 0.03 ng.L⁻¹. Satisfactory results were obtained for
206 field blank with Pt_D of 0.05 ng.L⁻¹.

207

208 *Platinum in SPM*

209 Particulate platinum concentrations (PtP) in Suspended Particulate Matter samples were
210 analyzed by Triple Quadrupole-Inductively Coupled Plasma-Mass Spectrometry (TQ-ICP-MS,
211 Thermo® iCAP TQ). Teflon filters with SPM (typical mass of ~ 3 - 5 mg) were acid digested
212 as described in Abdou et al. (2018). Briefly, 2 mL HCl and 1 mL HNO₃ (30 % HCl and 65 %
213 HNO₃ Suprapur, Merck®) were added to the samples in polypropylene (PP) tubes (DigiTUBEs,
214 SCP SCIENCE®) and closed with caps. Samples were placed in a Teflon-coated heating block,
215 and digested at 110 °C for 3 h. Cooled contents were then diluted in 10 mL Milli-Q water,
216 centrifuged at 4000 rpm for 10 min (20 °C) and filters were discarded. Analyses were
217 performed applying the standard internal addition method (as previously explained). After
218 obtaining the raw signal, Hafnium-oxide (HfO⁺) interferences were monitored using the ¹⁹³Ir
219 signal that is highly interfered by HfO⁺ (Djingova et al., 2003). Raw signal was thereafter
220 mathematically corrected using ¹⁹⁴Pt / ¹⁹⁵Pt natural ratio and assuming that HfO⁺ is the
221 only/dominant spectral interference on those selected isotopes. Analytical quality control was
222 performed using the BCR®-723 road dust (Institute for Reference Materials and
223 Measurements) and Jsd-2 sedimentary rocks (indicative value from Geological Survey of
224 Japan) as available CRMs for Pt concentrations. Analyses of these CRMs by TQ-ICP-MS gave
225 satisfactory recovery values of 87% and 101% respectively (n = 3) and precision expressed as

226 RSD of ~ 5 %. The detection limit (D.L.) for P_{tp} ($n = 10$) was estimated to 0.34 ng.g^{-1} for
227 typical sediment sample mass of 4 mg. Satisfactory results were obtained for field blank with
228 P_{tp} below detection limit.

229

230 *Complementary measurements*

231 Parallel to sampling, other master variables (i.e. temperature, dissolved oxygen, salinity, and
232 turbidity) as well as chlorophyll-a (Chl-a) concentrations were measured *in situ* using a
233 submersible probe (OS316*Plus*, Idronaut®), previously calibrated on-board the R/V MASO or
234 in an on-site laboratory on the CNR Platform. Salinity was measured using the Practical Salinity
235 Scale and Turbidity with Nephelometric Turbidity Units. Seawater pH and redox potential (Eh)
236 were determined with a Sentix® 41 probe (PROFILINE, WTW) and a PH-25 probe
237 (CRISON®), both previously calibrated using the manufacturer's specifications and supplied
238 reagents (NIST pH standards).

239 Dissolved organic carbon (DOC) was analyzed with an automated analyzer (TOC-L,
240 Shimadzu®) as described in Sharp et al. (1993). Accuracy of DOC measurements was tested
241 using reference material (Deep Sea Reference, DSR, University of Miami) providing recovery
242 results > 90 %. Analytical precision expressed as %RSD was 5 % at the 0.5 mg.L^{-1} range. The
243 D.L. ($n = 10$) was estimated to 0.06 mg.L^{-1} . Particulate organic carbon (POC) was analyzed
244 with a LECO® CS-125 after carbonate elimination as described elsewhere (e.g. Schäfer et al.,
245 2002). Analytical quality was checked by measuring CRMs (e.g. LECO 501–503). Accuracy
246 was within 5 % of the certified values and the analytical error generally < 5 % (% RSD).
247 Ammonium concentrations were determined by colorimetric procedure (visible-UV
248 spectrophotometer, Thermo®, Genesys 20, Aminot et al., 1997). Analytical error of the method
249 is < 5 % (% RSD), accuracy within 5 % of the reference values, and the D.L. of $18 \text{ } \mu\text{g.L}^{-1}$.

250 Satisfactory results were obtained for field blank values of DOC, POC and NH_4^+
251 ($\sim 0.2 \text{ mg.L}^{-1}$, 0.04 %, and $\sim 30 \text{ }\mu\text{g.L}^{-1}$ respectively).

252

253 2.3. Data treatment

254 The correlation matrix for the different parameters was produced using the software PAST®
255 applying the linear r (Pearson) parametric correlation coefficient. The statistical significance
256 (p-value) was set to 0.05. Surfer® software (Golden Software Inc., ver. 13) was used to produce
257 the distribution maps of the analytical results and the data interpolation was performed using
258 Minimum Curvature algorithm. Due to the complexity of the port area, a fault traces file of the
259 port shape (Fig. 1) and a spatial resolution of 5 m were used to perform the calculation gridding.
260 The interpolated variable limits were fixed to minimum/maximum of the measured data for the
261 Pt_D map; and from 0 until the maximum of the measured data for the Pt_P map. For each
262 interpolation map, the software returns the coefficient of multiple determination (R^2), which
263 represents the goodness of fit of the model. When the regression equation fits well to the data,
264 R^2 is large and close to 1.

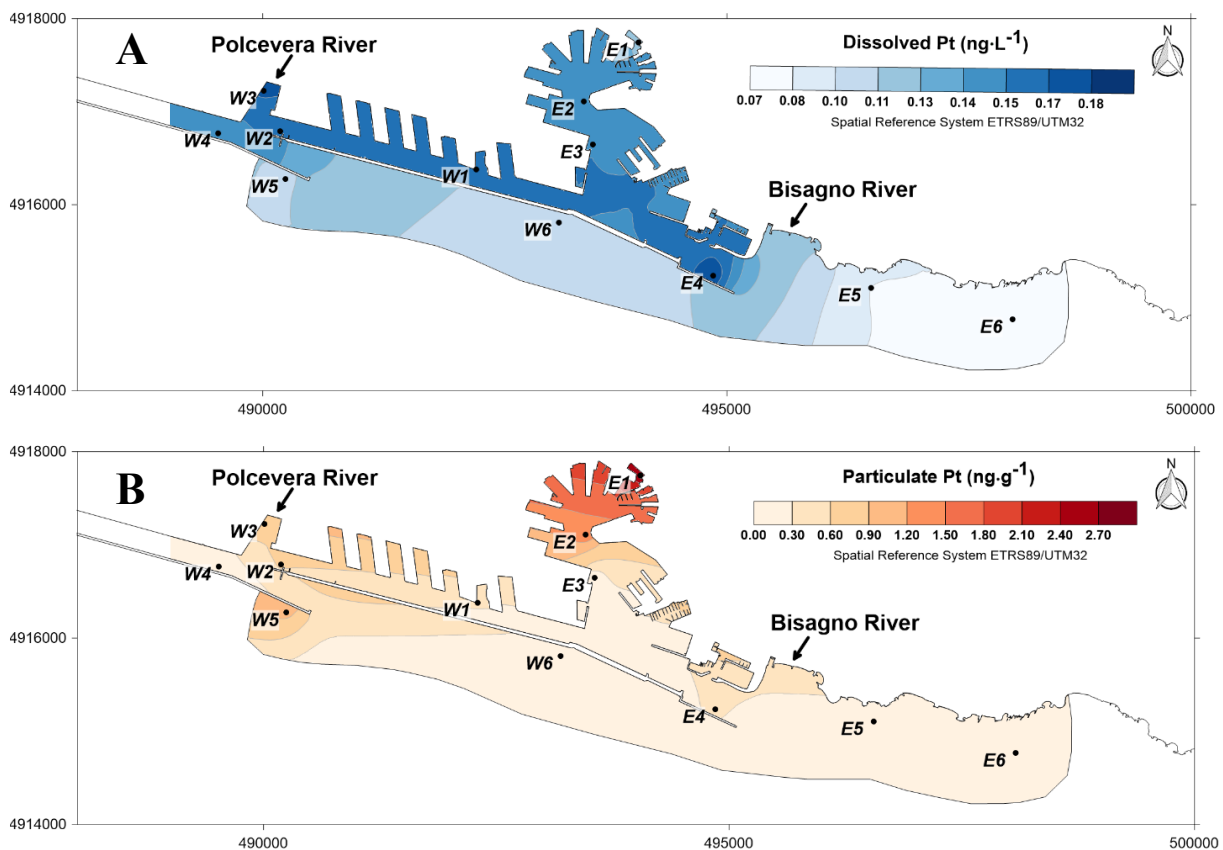
265 Particle–water distribution coefficients, K_D , defining the concentration ratios of Pt_P over Pt_D
266 were calculated according to the equation (1)

$$267 \quad K_D = [\text{Pt}_P] / [\text{Pt}_D] \times 10^3 \quad (1)$$

268 3. Results

269 The results obtained for major physical-chemical parameters and complementary
270 measurements are represented for two distinct longitudinal profiles, both starting from inside
271 the Genoa harbor and going towards the open Mediterranean Sea along the east and west part
272 of the coastline (Table 1). A clear decreasing gradient occurred for turbidity values from inside
273 towards the open sea eastward while westward profile featured no clear trend. Temperature
274 values varied between 14.5 °C and 15.0 °C with a trend for higher temperature for sampling
275 sites located inside the breakwaters. For both profiles, lowest temperatures occurred for external
276 sites being E6 and W6. For the eastern profile, salinity (S) of ~ 37.7 in the inner harbor,
277 compared to S = 38.0 in the outmost sites, suggested a ~ 1 % mixing of freshwater. Westward,
278 only the site W1 showed lower salinity (37.8), whereas other sites showed average salinity of
279 ~ 38. Redox potential (Eh) showed an increasing trend for eastward inner sites with highest
280 value of 211 mV for site E4, while at external sites E5 and E6, lower values (~ 190 mV)
281 occurred. Westward, Eh values did not show any particular trend (~ 180 mV), yet slightly
282 higher values at external sites W5 and W6 (~ 204 mV). Measurements of pH varied between
283 8.14 and 8.24, with highest values for external sites in both profiles (Table 1). Oxygen
284 saturation levels, DOC, and POC ranged between ~ 70 % and 98 %; 0.7 and 2.0 mg.L⁻¹; and
285 1 % to 5 % respectively. An increasing gradient occurred for O₂ saturation eastward from inner
286 to outer sites while levels of DOC, POC%, Chl-a and NH₄⁺ decreased (except for one higher
287 POC% value at site E5). Chlorophyll-a and NH₄⁺ values at sites E5 and E6 were of
288 ~ 0.3 µg.L⁻¹ and below the D.L. (18 µg.L⁻¹), respectively. In contrast, the westward profile
289 showed relatively constant values for O₂ saturation (~ 90 %). Elevated Chl-a, DOC, and POC %
290 levels tended to occur for stations inside the breakwaters, while most NH₄⁺ concentrations were
291 below the D.L.

292 Dissolved Pt concentrations ranged from 0.08 ng.L⁻¹ to 0.18 ng.L⁻¹ (Table 1) and showed clear
293 concentration differences between sites inside the harbor and sites outside the breakwaters
294 (Fig. 2A). Despite the distance from the stations, a coefficient of multiple determination of 0.51
295 was obtained for P_{TD} map that clearly confirms the observed gradient (Fig. 2A). Westward
296 profile showed somewhat similar P_{TD} concentrations at the sites W1 to W3, all inside the
297 breakwaters, and a decrease towards the most distant site of this profile (~ 0.10 ng.L⁻¹ at W6).
298 The eastward profile displayed an increasing P_{TD} gradient from ~ 0.10 ng.L⁻¹ at E1 to
299 0.18 ng.L⁻¹ at E4 and a sharp drop towards lower values at sites E5 and E6 (0.08 ng.L⁻¹;
300 Table 1).



301
302 **Fig. 2:** Dissolved and particulate Pt concentrations along the two longitudinal profiles in the
303 Genoa Harbor. A: spatial distribution of dissolved Pt concentrations (P_{TD}; ng.L⁻¹); and B: spatial
304 distribution of particulate Pt concentrations (P_{TP}; ng.g⁻¹).

305 The observed Pt_P concentrations ranged from below D.L. ($< 0.34 \text{ ng}\cdot\text{g}^{-1}$) up to $\sim 3 \text{ ng}\cdot\text{g}^{-1}$
306 (Table 1). A coefficient of multiple determination of 0.61 was obtained for the Pt_P map. Along
307 the westward profile, Pt_P concentrations were relatively constant and low ($< 1 \text{ ng}\cdot\text{g}^{-1}$; Fig. 2C)
308 with values below D.L. in sites W4 and W6. In the eastward profile, Pt_P were highest in the Old
309 Port (site E1), and showed an overall decreasing trend towards the outer sites reaching values
310 below D.L. at sites E5 and E6 (Pt_P level at site E3 was also below the D.L.).

311 Calculated particle–water distribution coefficients varied from 2×10^3 to 23×10^3 and showed
312 eastward and westward gradients similar to those observed for Pt_P concentration variations
313 (Table 1).

314 The relationships between spatial variations of Pt_D and Pt_P concentrations and the other
315 parameters measured are displayed separately for each profile only for the sites within the
316 breakwaters (Table 2). Despite the low number of samples, results of the correlation matrix
317 suggest that linear correlations exist between selected master variables and Pt levels in both the
318 dissolved and particulate phases along the profiles (Table 2). Eastward, Pt_D concentrations
319 seemed to be slightly positively correlated ($r \sim 0.6$) with wind speed and pH, and significantly
320 positively correlated ($r > 0.9$) with Eh and O_2 . These concentrations were significantly
321 negatively correlated with turbidity, DOC, POC Chl-a, and with NH_4^+ levels ($r < -0.9$).
322 Opposite behavior occurred for Pt_P concentrations. In the westward profile, Pt_D concentrations
323 which were weakly positively correlated with turbidity, Eh, ($r > 0.8$) and pH ($r \sim 0.6$) and
324 negatively correlated with $O_2\%$ ($r \sim -0.6$). Again Pt_P concentrations showed opposite behavior
325 with significant correlations (Table 2).

326 Platinum concentrations at the CNR Platform (fixed site PF; Fig. 1) measured at three different
327 times of the day showed variations for both Pt_D (0.09 to $0.15 \text{ ng}\cdot\text{L}^{-1}$) and Pt_P (3.1 to $14 \text{ ng}\cdot\text{g}^{-1}$;
328 Table 3), with highest concentrations at 12 pm. Dissolved Pt concentrations were in the same

329 range as those measured along the profiles, whereas relatively higher P_{tP} levels occurred,
330 leading to higher K_D values. During the measurements at the CNR Platform, wind direction
331 changed from north to south with relatively weak wind speed, similar to those during the
332 sampling of the westward profile. The master variables at the CNR Platform were in the same
333 range as levels observed at other internal sites (Tables 1 and 3). Only dissolved O_2 saturation
334 values were somewhat lower than those measured along the longitudinal profiles and were close
335 to those in the Old Port site (E1; Table 1). Dissolved Organic Carbon was also in the lower
336 range compared to longitudinal profiles, with values similar to those observed at site E4.
337 Ammonium values were close to those quantified during the westward profile.

338

339 **4. Discussion**

340 4.1. Water masses in the harbor and Pt distribution

341 Inside the harbor, water mass characteristics are supposedly strongly influenced by the
342 wind/wave conditions, the freshwater supply by the two main water courses (the Polcevera and
343 Bisagno Rivers), and run-off in the port area (Capello et al., 2010). Such physical parameters
344 may influence Pt distribution in the Genoa Harbor. Cutroneo et al. (2012) report the formation
345 of eddies in the current formed by wind in the Genoa area. The relative shallowness of the port
346 area (Table 1) further accentuates the influence of the wind on the entire water column (Capello
347 et al., 2010). Higher wind speed as recorded during the first day (Table 1) was correlated with
348 O_2 saturation and turbidity ($r \sim 0.8$ and -0.8 respectively; data not shown). This result suggests
349 an influence of wind speed on water mixing along the eastward profile. Along the westward
350 profile, weaker winds (coming from the south and probably stopped by breakwaters) showed
351 no correlation with O_2 saturation levels. Considering only the eastward profile, increasing wind
352 and subsequent water mixing is correlated with increasing P_{tD} and decreasing P_{tP} levels ($r \sim 0.6$

353 and -0.9 respectively). In the inner harbor, strong wind could lead to vertical and horizontal
354 seawater mixing, sediment resuspension, and redistribution of dissolved and especially
355 particulate components throughout the water column as observed in other studies (e.g. El Sayed
356 et al., 1994; see *Section 4.3.*).

357 The two profiles sampled westward and eastward from the Old Port show elevated Pt_D
358 concentrations within the whole harbor area, compared to the external sites (E5, E6 and W5,
359 W6). The lowest Pt_D values occurred at the sites E5 and E6 (Fig. 2A) mainly influenced by
360 open Mediterranean water transported with the east-west oriented northern Ligurian Current.
361 Therefore, the Pt_D concentrations in seawater from these sites ($\sim 0.08 \text{ ng.L}^{-1}$) may be considered
362 as representative of surface water in the northwestern Mediterranean Sea. This assumption is
363 supported by identical Pt_D levels of 0.08 ng.L^{-1} in surface waters from an open-sea site, south
364 from the Balearic Islands (Abdou, 2018). Open Mediterranean surface water Pt_D concentrations
365 appear somewhat higher than average Pt_D in the North Atlantic seawater reported in the early
366 nineties ($\sim 0.05 \text{ ng.L}^{-1}$; Colodner, 1991) and recently confirmed by López-Sánchez et al. (2019)
367 for North, Central and South West Atlantic Ocean. Values are also higher than those reported
368 in the northwest Pacific Ocean ($\sim 0.04 \text{ ng.L}^{-1}$; Suzuki et al., 2014). However, the supposed
369 Mediterranean surface water Pt_D is similar to values in the coastal North Atlantic Ocean off the
370 slightly Pt-contaminated Gironde Estuary (0.08 ng.L^{-1} ; Cobelo-García et al., 2014a) and lies
371 within the range of values reported for the Lérez Estuary seawater end-member (from 0.08 to
372 0.10 g.L^{-1} ; Cobelo-García et al., 2013). These observations are consistent with expected
373 relatively higher Pt inputs into the semi-enclosed Mediterranean Sea due to the strong
374 urbanization of its coastline. Similarly, the western profile shows the lowest Pt_D values (i.e.
375 $\sim 0.10 \text{ ng.L}^{-1}$; Fig. 2A) at the sites located outside the breakwaters (W5 and W6). These
376 concentrations are slightly higher than open Mediterranean Pt_D , suggesting minor local
377 contamination off the harbor and ‘downstream’ from the sites E5 and E6 (with respect to the

378 dominant westward Ligurian Current). Conversely, inside the breakwaters, contamination is
379 consistently higher with Pt_D concentrations up to more than 2-fold the open Mediterranean
380 value (Fig. 2A).

381 Particulate Pt concentrations in suspended particles partly reflect the general concentration
382 gradient from inside the harbor towards the open sea. They were measurable (~ 0.4 to 3 ng.g^{-1})
383 at all internal sites and behind the breakwaters (except site E3), whereas for the external sites
384 from outside the breakwaters Pt_P were below D.L. ($< 0.34 \text{ ng.g}^{-1}$; Fig. 2B). The Pt_P levels inside
385 the Genoa Harbor were similar to those reported for the Lérez Estuary (~ 1 to 15 ng.g^{-1}),
386 interpreted as anthropogenic Pt enrichment (Cobelo-García et al., 2013). Such enrichment may
387 partly originate from small watercourses flowing into the harbor (Fig. 1), even though salinity
388 measurements show little influence of freshwater inputs. Since sampling took place at $\sim 5 \text{ m}$
389 depth i.e. below the main influence of freshwater torrents (Capello et al., 2016) and no strong
390 rainfall occurred the previous days (only 3.8 mm on April 2nd), our salinity records mainly
391 reflect seawater influence. In the inner part of the Old Port, the semi closed shape and shallow
392 bottom ($< 10 \text{ m}$ depth; Table 1) of the harbor make the water exchange times rather long, in
393 particular close to the bottom depth (corresponding to E1 sampling point). Povero et al. (2005)
394 calculated a mean of 20 days for a complete water exchange in this inner part and 7 days in the
395 outer system (central basin and eastern entrance), which is wider and $\sim 10 \text{ m}$ deeper. Although
396 both calculated water residence times are not representative of the whole area (because they do
397 not consider its irregular shape) and refer mainly to the eastward transect, they can help
398 explaining the Pt enrichment in the inner part of the harbor. They may also explain the
399 decreasing gradient towards the outer basin and the open sea, together with dilution affecting
400 the generally stagnant harbor waters when flowing outside. Moreover, inside the harbor,
401 important dredging activities were undertaken from 2009 to 2014 and are, to a lesser extent,
402 still carried out in the recent years. Since then, turbid plumes, influenced by wind/wave

403 conditions, have been recorded (Cutroneo et al., 2012). Intensification of big ships and ferry
404 transit and berthing increase the presence of turbid plumes (Capello et al., 2010), suggesting
405 that sediment resuspension may represent a source of trace metals, including Pt, in the system.

406

407 4.2. Anthropogenic Pt sources

408 *Industrial activities*

409 Along the eastern profile, from the inner to the outer part of the harbor, Pt_D concentrations
410 tended to increase with a maximum at site E4, close to the mouth of the Bisagno River (Fig. 2A),
411 while the highest Pt_D in the westward profile occurred inside the breakwaters, including site
412 W3 (0.17 ng.L⁻¹), i.e. near the Polcevera River mouth. The western part of the harbor hosts coal
413 deposits and a terminal for oil products (Fig. 1). Indeed, westward the Polcevera River mouth,
414 the Multedo oil terminal is one of the most important in Italy and Europe. It handles crude oil,
415 final products (petrol, gasoline, fuel oil) semi-manufactured, and basic petrochemical products,
416 for a maximum total capacity of 30 million tons per year (Recchia et al., 1999). Platinum is
417 commonly used as a catalytic agent in oil industry (e.g. catalytic cracking of crude petroleum;
418 Peavy, 1958) and such activities could therefore be a source of Pt inputs into the coastal system.
419 Built in 1952, the coal power plant of Genoa is also located on the west side of the harbor. Coal
420 industry may contribute to local Pt concentrations, since high Pt concentrations have been
421 reported for coal samples (~ 5 ng.g⁻¹, Finkelman and Aruscavage, 1981; up to 70 ng.g⁻¹, Dai et
422 al., 2003). The bulk terminal of the harbor containing coal (dry bulk) and oil (liquid bulk) is
423 also located in the western area (Cutroneo et al., 2017). Historical intense oil and coal-based
424 industrial activities in the Genoa Harbor and erosion/leaching of such deposits stored uncovered
425 against wind and/or rain might lead to the presence of industrial coal particles in the suspended
426 particulate matter around the Polcevera River mouth and in the Sampierdarena channel that

427 connects the outer basin of the Old Port and Polcevera River mouth. Although the Pt
428 concentrations in this oil or coal are unknown, one cannot exclude that oil inputs, industrial coal
429 and/or fly ashes could contribute to the Pt enrichment at this sampling site given the range of
430 values reported in the literature. Since 1953, a steel mill has also been present in the western
431 area and related activities cause the release of particulate matter of different sizes (Mazzei et
432 al., 2006). Metallurgic activities can be a source of Pt release into aquatic environments as
433 recorded in other systems (e.g. Gironde fluvial estuarine continuum; Abdou et al., 2016).
434 Furthermore, in urban systems, other possible Pt sources may arise from direct inputs of dental
435 laboratories, electronic industries, glass manufacturing, Pt-containing pharmaceuticals or
436 industrial catalyst productions (Kümmerer et al., 1999).

437

438 *Hospital and sewage effluents*

439 Platinum-based compounds have been successfully applied in anti-cancer chemotherapy for
440 decades (e.g. Lenz et al., 2007). Surface waters draining effluents from hospital with oncology
441 ward contain patient excretion of Pt-based anticancer drugs and display Pt concentrations that
442 could exceed natural inputs (Vyas et al., 2014). Monteiro et al. (2017) evaluated Pt_D
443 concentrations in the inflow and outflow of a wastewater treatment plant (WWTP) in Lisbon to
444 ~ 5 and 15 ng.L^{-1} respectively after a dry weather period and ~ 10 and 25 ng.L^{-1} after a raining
445 event. In the western part of the Genoa Harbor, Pt inputs from sewage treatment plants reach
446 the Polcevera river mouth (Fig. 1). Despite probable dilution in natural waters compared to
447 levels previously reported, such inputs may add to Pt sources affecting the sampling site W3
448 ($Pt_D \sim 0.18 \text{ ng.L}^{-1}$). Small sewage effluents also occur directly in the Old Port (Fig. 1).
449 Considering only the sites E1 to E4 located inside the relatively stagnant harbor waterbody (due
450 to confinement and low tidal amplitudes; Cutroneo et al., 2017, 2012), the eastward gradients
451 of Pt concentrations have been correlated with parameters reflecting sewage effluent signature.

452 Inside the harbor, Pt_D concentrations were positively correlated with dissolved oxygen ($r > 0.9$)
453 and pH ($r > 0.6$), and negatively correlated with turbidity, Chl-a, % POC, DOC, and NH_4^+
454 ($r < -0.9$). The opposite trends occurred for Pt_P concentrations, which were positively correlated
455 with Chl-a, % POC, DOC, and NH_4^+ . Assuming that these major variables serve as proxies for
456 wastewater inputs and degradation (Ruggieri et al., 2011), this could explain Pt distribution
457 inside the Old Port. Accordingly, supply of discharged sewage rich in organic matter and poor
458 in oxygen content typically occur into the harbor basins (Ruggieri et al., 2011). The CNR
459 platform (site PF) and surrounding waters are close to the urban area hosting a relevant center
460 for Medical Oncology (red cross/blue circle). Highest levels of Pt_P (14 ng.g^{-1}) are recorded at
461 this sampling site and relatively higher levels of Pt_D (0.15 ng.L^{-1}) both at 12 pm. For the same
462 sample, sewage-related signals are recorded, with clearly lower O_2 saturation and highest POC
463 levels compared to the other sites. Sites E1 and PF are also characterized by a more confined
464 area with shallower bottom depth ($\sim 6 \text{ m}$ depth). High levels of Pt_P and relatively low levels of
465 Pt_D , where sewage inputs are at maximum (site E1) might be related to: (i) the presence of
466 originally high Pt_P concentrations in wastewaters or run-off, and/or (ii) the adsorption of Pt_D on
467 organic-rich ligands present in wastewaters (high concentrations of dissolved and particulate
468 organic carbon in the turbid sewage plume). The effect of sediment resuspension in these
469 shallower sampling sites is not discarded. Kinetics of suspected adsorption/desorption
470 processes may explain the observed differences in Pt_D and Pt_P distributions between sites (see
471 *Section 4.3.*).

472

473 *Traffic-related emissions*

474 The Genoa Harbor is in direct vicinity ($< 5 \text{ km}$ distance) from the urban highway and very close
475 to urban heavy traffic roads, a likely source for Pt bearing particles. Automobile catalysts,
476 which are thought to be the principal Pt anthropogenic source, have atmospheric residence times

477 ranging from hours to days depending on particle size and density and can be transported from
478 regional to longer distances (Rauch et al., 2006, 2004). Dust and suspended matter with Pt
479 particles are thereafter removed from the atmosphere by wet (rainfall) or dry (gravitation)
480 deposition (Pawlak et al., 2014). For other trace metals such as copper (Cu), enhanced
481 atmospheric input of Cu-rich urban particles showing great water solubility may result in the
482 predominance of the atmospheric input of Cu over the river input (El Sayed et al., 1994).
483 Therefore, in addition to causing water mixing, wind can be a vector of Pt into aquatic systems.
484 The two wind situations observed during our study are typical of the Genoa area. Winds from
485 the N-NE (0–30° N) and SE (135–150° N) are two of the most frequent wind situations in
486 winter and summer, respectively (Cutroneo et al., 2012). Our field campaign took place in
487 spring, a period of the year generally featuring the two situations. The eastward profile was
488 sampled under relatively strong N-NE winds causing the currents in the entire harbor to flow
489 outwards (Cutroneo et al., 2012). Relatively clear gradients of all monitored parameters can be
490 explained by such conditions. Recent data confirm the importance of atmospheric deposition in
491 Pt enrichment of oceanic surface waters (López-Sánchez et al., 2019). Automobile catalyst
492 exhaust Pt inputs come from the land and Pt-bearing particles can be transported by the wind
493 causing a gradient of Pt_P concentrations eastward. In the Genoa Harbor system, such particles
494 could originate from catalyst exhaust fumes and/or industrial fly ashes. On the reverse, the
495 westward profile was carried out under weaker, SE winds and showed less clear parameter
496 variations. The E and SE winds generate a westward current forcing the surface harbor waters
497 inside, while only in the bottom layer a weak outflow can occur (Cutroneo et al. 2017, 2012).
498 However, due to the morphological complexity and the port structures jutting out in all
499 directions, there is an extreme variability in the current direction in the port (Capello et al.,
500 2010) and more data would be needed covering longer time-scales to confirm the present
501 assumptions. If such source types influence Pt partitioning and its spatial distribution, they also

502 may affect Pt levels both in historical records (e.g. Abdou et al., 2019, 2016) and at short
503 temporal scales as observed in site PF.

504

505 4.3. Platinum reactivity and partitioning

506 *Platinum seawater speciation*

507 Platinum concentrations in both dissolved and particulate phases are controlled by several
508 factors implying various biogeochemical processes. Assuming that Pt originates from multiple
509 potential point sources prevailing at different sites it is likely that Pt speciation differs from site
510 to site. Organic forms might be represented by the three Pt-based drugs mainly used in the
511 hospital: cisplatin, carboplatin and oxaliplatin along with their metabolites, while Pt emitted
512 from vehicles or industrial activities may be under inorganic forms. Hospital effluents transport
513 dissolved Pt ranging from $< 10 \text{ ng.L}^{-1}$ to $250 \text{ }\mu\text{g.L}^{-1}$ (Rauch and Peucker-Ehrenbrink, 2015;
514 Vyas et al., 2014), mainly excreted from these anticancer drugs and/or derived metabolites.
515 Such excretion also continues outside the hospital, as more and more patients go home directly
516 after the treatment to the extent that the majority of Pt is emitted from outpatients at home (Lenz
517 et al., 2007; Vyas et., 2014). The phase in which Pt species are released as well as the
518 dissolved/particulate partitioning in sewage and the related removal efficiency in WWTPs
519 remain widely unknown. Previous studies suppose that municipal and hospital effluents contain
520 carboplatin (the most stable drug) as well as degradation products being the monoaquacisplatin
521 and the diaquacisplatin, from anti-cancer drug administration (Lenz et al., 2007; Vyas et al.,
522 2014). Recent work reports that the behavior of organic Pt species in recipient waters is
523 governed by chloride (Cl^-) concentrations with, for cisplatin, decreasing proportion of reactive
524 species with increasing salinity (Curtis et al., 2010; Turner and Mascorda, 2015). Curtis et al.
525 (2010) proved that environmental conditions favoring cisplatin retention and its adsorption on

526 particles are turbid waters of low chlorinity and long residence time. Dispersal is likely to occur
527 in rapidly flushed waters under low turbidity conditions and high chlorinity. Our study site is
528 characterized by high salinities that may limit cisplatin adsorption, but turbid plumes arriving
529 from sewage effluents and long water and particle residence times inside the breakwaters may
530 in turn favor such process. More investigation is needed under similar environmental
531 conditions.

532 Both run-off and sewage also contain PGE-bearing catalytic converter fragments with intact
533 PGE-load (Prichard and Fisher, 2012), which may undergo partial dissolution in contact with
534 seawater. The few data on Pt speciation in seawater suggest that Pt(IV) is the most important
535 oxidation state (Cobelo-García et al., 2013). Laboratory works showed that its inorganic
536 equilibrium speciation is dominated by the stable $\text{PtCl}_5(\text{OH})^{2-}$ complex (Gammons, 1996;
537 Cobelo-García et al., 2014a). Laboratory and field observations report decreasing particulate-
538 aqueous distribution coefficient for inorganic forms with increasing salinity (Cobelo-García et
539 al., 2014, 2008; Turner, 2007), suggesting increasing Pt levels in the dissolved phase as Cl^-
540 concentrations increase. The observed eastward particulate/dissolved gradients suggest Pt
541 dissolution from solid carrier phases, either road dust (Sebek et al., 2011) or small sewage
542 particles, and subsequent dilution of Pt_D by seawater that would be favored by Pt-Cl complex
543 formation.

544 *Potential carrier phases of Pt in sites with anthropogenic influence*

545 Given the above, one would expect inputs of different co-existing Pt forms from different
546 overlapping sources into the marine environment with wastewater, run-off of the surrounding
547 area and atmospheric deposition. Numerous discharges occur in the Genoa Harbor area,
548 comprising release of unknown and potentially dangerous substances (Fig. 1) that may
549 contribute to Pt contamination. A rough mass balance ($Pt_D = Pt_P \times SPM \times 10^{-3}$) for the eastern
550 profile inside the harbor suggests that Pt release from suspended particles would account for a
551 Pt_D addition of $\sim 0.02 \text{ ng.L}^{-1}$, whereas the observed Pt_D values increased by $\sim 0.06 \text{ ng.L}^{-1}$. The
552 amount of Pt presumably transferred from the particulate to the dissolved phase is not
553 contradictory to the field data but does not explain all the Pt_D addition observed, supporting the
554 idea of additional Pt_D inputs through the different discharges (Fig. 1). Outside the breakwaters,
555 biogeochemical equilibration processes and the resulting higher Pt_D would be masked by
556 massive dilution with open Mediterranean water. Biogeochemical processes, inherent to
557 anthropogenic discharges and the mixing of environmental waters can widely modify the
558 original Pt phase. Previous work has shown that relatively high dissolved metal loads in
559 wastewater inputs equilibrate in the recipient waters. Efficient adsorption on high complexation
560 site particles induces profound changes in element partitioning between the dissolved and
561 particulate phases (e.g. Deycard et al., 2014). Indeed, the residual POC in effluent wastewater
562 mainly consists of reactive (degradable) organic matter, once exposed to natural microorganism
563 communities in the receiving waterbody (Deycard et al., 2017). Such organic matter and
564 especially that containing thiols and reduced sulfur groups, may strongly bind soft metals and
565 potentially compete with inorganic ligands such as Cl^- from the recipient waters (Levard et al.,
566 2013; Wood et al., 1990). Literature suggests that, considering cisplatin flocculation in natural
567 water, complexation with specific organic ligands containing nitrogen or sulphur is likely (e.g.
568 Curtis et al., 2010; Turner and Mascorda, 2015). This means that Pt partitioning between the

569 two phases might be partly influenced by interactions with POC from wastewater discharges
570 when mixing with coastal waters, although the degree of surface complexation with organic
571 matter remains unknown (Couceiro et al., 2007). Natural sediments may also contain POC and
572 laboratory experiments have proven that Pt adsorption kinetics onto estuarine sediments is
573 greatest for cohesive sediments with relatively high total carbon contents (Couceiro et al.,
574 2007). Another experimental study showed that complexation with organic ligands can be
575 critical to particle-water fractionation of Pt discharged into aquatic systems, removal from the
576 dissolved phase being most favorable when entering a turbid watercourse (Turner et al., 2006).
577 All of these observations seem in line with the reported highest Pt_P levels co-occurring with
578 high turbidity sewage inputs in the Genoa Harbor (Tables 1 and 3; Fig. 2B). Depth profiles of
579 Pt_D concentrations in the Atlantic Ocean also showed a higher particle-reactivity or lower
580 solubility of Pt_D with decreasing oxygen levels (López-Sánchez et al., 2019).

581 The related daily variations in K_D values at site PF (Table 3) suggest that the observed K_D might
582 not describe equilibrium conditions. However, the average K_D values obtained for the Genoa
583 Harbor ($\sim 8 \times 10^3$) are consistent with values obtained for the seawater end-member of the
584 Gironde Estuary ($\sim 4 \times 10^3$, Cobelo-García et al., 2014a), and the maximum K_D value recorded
585 at the CNR Platform ($\sim 1 \times 10^5$) is similar to those found in the seawater end-member of the
586 Lérez Estuary (Cobelo-García et al., 2013). Outside the Old Port, but still inside the harbor, the
587 growing influence of open seawater with a decrease in turbidity, as well as change in the nature
588 of SPM surfaces (decrease in %POC) could explain low Pt_P concentrations (below detection
589 limit), going along with increasing Pt_D . Considering possible wind effect and water column
590 mixing, remobilization of Pt from particles through the degradation of organic matter and redox
591 reactions (correlation of Pt levels with pH and Eh parameters; Table 2) seems to contribute to
592 Pt_D enrichment at those sampling sites, as observed elsewhere for Pt and other trace metals
593 (Almécija et al., 2016; El Sayed et al., 1994). However, multiple solid phases other than solely

594 POC can also decrease Pt_D concentrations. Inorganic mineral phases such as iron and
595 manganese oxides may also be important Pt carrier phases (Colodner et al., 1992). In fact, those
596 phases could be more important than organic material for the removal of cisplatin degradation
597 products (Turner and Mascorda, 2015).

598 Whatever the dissolved or particulate phase transporting Pt, this element has proven to be
599 bioavailable to marine organisms (e.g. Rauch et al., 1999). It is crucial to better understand Pt
600 speciation and distribution, since the physical and chemical forms of this emerging contaminant
601 are expected to considerably influence its biological availability. In fact, previous work on Pt
602 toxicity in marine bivalves has shown strong accumulation and the onset of defense
603 mechanisms, for exposure levels that might be reached in case of acute pollution, especially in
604 semi-closed coastal systems (Abdou et al., 2018).

605

606 4.4. Coastal anthropogenic contamination – Pt as a novel urban tracer

607 Although the geochemical cycles of Pt in coastal environments under high urban/industrial
608 pressure are far from being fully understood (due to the limited number of data), our study,
609 combined to already existing works, allows to draw perspectives for future research in this field.
610 Assuming the dominance of Pt organic forms released by hospital effluents, monitoring of Pt
611 concentrations may serve as a tracer for these effluents release and spreading in the environment
612 together with other proxies commonly used such as POC% or nutrient levels. Main substances
613 expected in effluents are the most stable anti-cancer compounds (including carboplatin) and
614 degradation products from the different drugs (Lenz et al., 2007; Vyas et al., 2014). However,
615 important behavior differences (reactivity, phase affinity, removal...) are observed among Pt-
616 based drugs and their various degradation products (Turner and Mascorda, 2015). Such organic
617 Pt sources need monitoring and characterization, especially considering the increasing cancer

618 incidence and the related increasing demand for chemotherapy treatment in developed countries
619 (Johnson et al., 2008). Inorganic forms of Pt are assumed to dominate in catalytic converters
620 emissions. High resolution monitoring of concentrations of those forms could therefore reflect
621 atmospheric particle deposition and traffic-related urban contamination in coastal areas.
622 Inorganic Pt forms may also reflect atmospheric contamination related to industrial activities
623 including oil and coal-based industries. The lack of knowledge regarding Pt speciation is
624 directly related to analytical challenges inherent to the ultra-trace environmental levels of this
625 emerging metallic contaminant. Future work should focus on developing analytical techniques
626 for routine Pt quantification in seawater for both dissolved and particulate phases including
627 speciation analyses.

628 Spatial distribution of Pt concentrations in the Genoa Harbor showed disparities partly
629 depending on urban source proximity, yet, temporal aspects also may be crucial. Dredging
630 activities, daily cycles in traffic intensity, industrial emissions, and/or wastewater composition
631 (Deycard et al., 2014; Mazzei et al., 2006; Kümmerer and Helmers, 1997), as well as wind
632 conditions could lead to variable fluxes of Pt into the harbor waters explaining why Pt_P varied
633 by a factor of ~ 5 (i.e. from 3 to 14 $ng.g^{-1}$ at site PF) within few hours. Spatial and temporal
634 variations in Pt levels inside the Genoa Harbor may therefore reflect both distance from the
635 source and release dynamics of a wide range of different potential sources.

636 Platinum distribution therefore appears to vary according to urban pollution type, source and
637 intensity, suggesting that this element can serve as an innovative and sensitive tracer of
638 anthropogenic impact from multiple urban sources (traffic, sewage, hospital effluents). The
639 present dataset suggests that such anthropogenic inputs supplied to rivers or directly in the
640 harbor by urban and/or industrial wastewater discharges and by atmospheric deposition may
641 represent important local Pt point sources to coastal waters. Platinum levels observed in the
642 Genoa Harbor waters are far lower than concentrations measured in highly contaminated waters

643 including polluted rivers (up to $\sim 7 \text{ ng.L}^{-1}$ in Tokyo Bay urban rivers; Obata et al., 2006) or
644 even hospital effluents (up to $250 \text{ }\mu\text{g.L}^{-1}$; Lenz et al., 2007; Vyas et al., 2014). It remains unclear
645 to what extent dilution of Pt concentrations by recipient seawater or the absence of rainy events
646 during our sample collection could have resulted in relatively low Pt levels. Future research and
647 field sampling should therefore consider sample collection both near pollution sources (i.e.
648 rivers, urban discharges, sewage effluents) and in recipient coastal waters in order to perform
649 relevant mass balance calculations. This should be done at high spatial as well as high temporal
650 resolutions in order to fully understand Pt cycles in urban coastal environments.

651 **5. Conclusions**

652 With TCE increasing environmental release being a global concern, this work provides
653 knowledge on the extent of Pt contamination and biogeochemical processes occurring in a
654 representative urban coastal site. The first information on Pt distribution and reactivity of both
655 dissolved and particulate phases in an industrialized and urbanized Mediterranean harbor is
656 given. Measured concentrations for Pt_D (~ 0.08 to 0.18 $ng.L^{-1}$) and Pt_P (< 0.34 to 14 $ng.g^{-1}$)
657 allowed proposing $Pt_D \sim 0.08$ $ng.L^{-1}$ (as measured in the Ligurian Current and south from the
658 Balearic Islands) as a first reference value for western Mediterranean surface seawater. There
659 is a clear evidence of anthropogenic Pt contamination at all sites studied within the Genoa
660 Harbor showing distinct concentration gradients. They are attributed to different overlapping
661 local sources such as industry, run-off, hospital effluents, domestic wastewater as well as
662 physical parameters such as wind and dust / catalyst particles atmospheric deposition. Gradients
663 in both Pt_D and Pt_P as well as major variables attributed to input and degradation of urban
664 sewage inside the relatively stagnant harbor waters reflect biogeochemical processes modifying
665 the nature, surface properties and abundance of suspended particles, leading to profound
666 changes in Pt partitioning and carrier phases. While quantification of Pt emitted from vehicle
667 exhaust, supposedly under inorganic form, rise attention, organic forms may also be of major
668 concern in urbanized coastal systems. Such mobilization of anthropogenic Pt can enhance its
669 bioavailability, bio-uptake and induce adverse effects on marine organisms. The role of
670 biogeochemical processes and responses from marine wildlife to contamination, including
671 potential Pt complexation to biologically-derived organic ligands such as algae exudates, are
672 still unknown and need further investigation. On the other hand, this study opens the perspective
673 of using Pt as an innovative tracer of urban inputs that should be applied in other urbanized
674 coastal areas of the Mediterranean Sea and worldwide.

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682

683 **Conflicts of interest**

684 The authors declare no conflicts of interest.

685

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911

912 **Tables**

913 **Table 1:** Dissolved and particulate Pt concentrations, Pt distribution coefficients, and master
 914 variables monitored along two longitudinal profiles in the Genoa Harbor. The eastward profile
 915 corresponds to sites E1 to E6, with E1 being located in the Old Port, while the westward profile
 916 corresponds to sites W1 to W6. With: Dissolved Pt concentration (Pt_D ; $ng.L^{-1}$), Particulate Pt
 917 concentration (Pt_P ; $ng.g^{-1}$), Particle-water distribution coefficient (K_D), Wind direction (Wind
 918 dir.), Wind speed (Wind sp.; $km.h^{-1}$), Bottom depth (m), Turbidity (Turb.; NTU), Temperature
 919 (Temp; $^{\circ}C$), Salinity, Redox potential (Eh; mV), pH, dissolved oxygen saturation ($O_2\%$),
 920 Dissolved Organic Carbon (DOC; mg/L), Particulate Organic Carbon (POC%), Chlorophyll-a
 921 (Chl-a; $\mu g/L$), ammonium (NH_4^+ ; $\mu g/L$), d.l.: detection limits, n.d.: not determined.

Param./Site	W6	W5	W4	W3	W2	W1	E1	E2	E3	E4	E5	E6
Pt_D	0.10	0.11	0.14	0.17	0.16	0.17	0.12	0.15	0.15	0.18	0.08	0.08
Pt_P	<d.l.	0.99	<d.l.	0.65	0.65	0.37	2.6	1.4	<d.l.	0.50	<d.l.	<d.l.
K_D ($\times 10^3$)	n.d.	9.1	n.d.	3.7	4.1	2.2	23	9.9	n.d.	2.8	n.d.	n.d.
Wind dir.	ESE	SSE	SSE	SE	S	SSE	N	NNE	N	NE	N	N
Wind sp.	15	9	7	9	6	6	7	17	22	15	22	15
Bottom depth	24.3	27.6	11.0	13.0	11.0	9.6	6.0	9.7	10.7	16.0	12.5	15.0
Turb.	2.23	7.38	2.89	5.17	2.51	4.17	6.47	4.85	3.88	3.52	2.10	2.13
Temp.	14.61	14.82	14.88	14.80	14.84	15.02	14.94	14.84	14.85	14.96	14.50	14.88
Salinity	38.0	37.9	38.0	38.0	38.0	37.8	37.8	37.7	37.8	37.8	38.1	38.0
Eh	204	203	168	187	187	179	173	182	196	211	181	196
pH	8.24	8.21	8.17	8.22	8.22	8.18	8.14	8.14	8.19	8.17	8.21	8.20
$O_2\%$	94.5	92.5	93.9	90.4	96.0	92.6	69.1	87.9	91.4	94.0	97.9	96.8
DOC	1.05	1.17	1.16	0.98	0.84	1.14	2.05	1.28	1.10	0.68	0.91	1.04
POC%	3.75	2.81	2.60	4.87	2.32	0.89	4.81	2.73	3.11	2.47	4.35	1.94
Chl-a	0.37	1.02	1.12	1.48	0.59	0.95	1.16	0.98	0.82	0.82	0.25	0.35
NH_4^+	<d.l.	25.1	<d.l.	41.5	<d.l.	<d.l.	137	113	111	102	<d.l.	<d.l.

922

923 **Table 2:** Correlation table of Pt_D and Pt_P concentrations with master variables. Values correspond to
 924 Pearson's r coefficient. n.d.: not determined. Stars correspond to a significant level of 0.05 (p -
 925 value<0.05). Note that only the sites inside the breakwaters are considered.

	Eastward profile (E1 to E4)		Westward profile (W1 to W4)	
	Pt_D	Pt_P	Pt_D	Pt_P
Wind sp.	0.59	-0.90	0.43	0.52
Turb.	-0.95*	1.00*	0.80	-0.12
Temp.	0.15	-0.14	-0.07	-0.99*
Salinity	0.43	-0.34	-0.39	0.99*
Eh	0.96*	-0.93	0.83	1.00*
pH	0.63	-0.86	0.63	1.00*
O₂%	0.92*	-0.98*	-0.62	0.10
DOC	-0.99*	1.00*	-0.35	-0.87
POC%	-0.90*	0.94	0.36	0.79
Chl-a	-0.91*	1.00*	0.33	0.12
NH₄⁺	-0.97*	0.99*	n.d.	n.d.

926

927 **Table 3:** Dissolved and particulate Pt concentrations, particle-water distribution coefficients,
 928 and master variables at the CNR Platform (site PF) at 10 am, 12 pm, and 2 pm. Units of the
 929 various parameters are as reported in Table 1.

Param./Time	10 am	12 pm	2 pm
Pt_D	0.09	0.15	0.11
Pt_P	7.7	14	3.1
K_D ($\times 10^3$)	82	94	27
Wind dir.	N	SSE	S
Wind sp.	4	9	7
Bottom depth	6	6	6
Turb.	4.00	2.60	4.60
Temp.	14.76	14.66	14.73
Salinity	38.0	38.0	38.0
Eh	190	182	191
pH	8.20	8.20	8.21
O₂%	81.9	82.8	87.1
DOC	1.01	0.68	0.63
POC%	2.50	5.15	2.02
Chl-a	1.00	1.40	1.60
NH₄⁺	47.0	48.1	44.8

930

931 **Figure captions**

932 **Fig. 1:** Sampling sites in the Genoa Harbor. Two sampling profiles were performed with an
933 eastward profile (sites E1 to E6) and a westward profile (sites W1 to W6). Sampling site PF
934 (PlatForm) corresponds to the CNR Platform. White arrows indicate river mouths. WWTPs:
935 Wastewater Treatment Plants.

936 **Fig. 2:** Dissolved and particulate Pt concentrations along the two longitudinal profiles in the
937 Genoa Harbor. A: spatial distribution of dissolved Pt concentrations (Pt_D ; $ng.L^{-1}$); and B: spatial
938 distribution of particulate Pt concentrations (Pt_P ; $ng.g^{-1}$).

939

940 **List of tables:**

941 **Table 1:** Dissolved and particulate Pt concentrations, Pt distribution coefficients, and master
942 variables monitored along two longitudinal profiles in the Genoa Harbor. The eastward profile
943 corresponds to sites E1 to E6, with E1 being located in the Old Port, while the westward profile
944 corresponds to sites W1 to W6. With: Dissolved Pt concentration (Pt_D ; $ng.L^{-1}$), Particulate Pt
945 concentration (Pt_P ; $ng.g^{-1}$), Particle-water distribution coefficient (K_D), Wind direction (Wind
946 dir.), Wind speed (Wind sp.; $km.h^{-1}$), Bottom depth (m), Turbidity (Turb.; NTU), Temperature
947 (Temp; $^{\circ}C$), Salinity, Redox potential (Eh; mV), pH, dissolved oxygen saturation ($O_2\%$),
948 Dissolved Organic Carbon (DOC; mg/L), Particulate Organic Carbon (POC%), Chlorophyll-a
949 (Chl-a; $\mu g/L$), ammonium (NH_4^+ ; $\mu g/L$), d.l.: detection limits, n.d.: not determined.

950 **Table 2:** Correlation table of Pt_D and Pt_P concentrations with master variables. Values
951 correspond to Pearson's r coefficient. n.d.: not determined. Stars correspond to a significant
952 level of 0.05 ($p\text{-value}<0.05$). Note that only the sites inside the breakwaterss are considered.

953 **Table 3:** Dissolved and particulate Pt concentrations, particle-water distribution coefficients,
954 and master variables at the CNR Platform (site PF) at 10 am, 12 pm, and 2 pm. Turb.: turbidity.
955 Units of the various parameters are as reported in Table 1.