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Title: VARIABLES AFFECTING THE PLANKTON NETWORK IN MEDITERRANEAN PORTS

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Abstract: Attention on port waters is increasing since these economically important infrastructures are embedded in the coastal environment and their management needs to be considered in the monitoring programmes of coastal ecosystems. To implement the sustainable development (blue growth) of port areas, a general knowledge on the ongoing processes in their waters needs to be obtained, considering both abiotic and biotic variables. The present study aimed at inspecting the relationships among plankton components to provide insights into ecology of ports. Seasonal samplings were carried out in three Mediterranean touristic ports where bacterio-, phyto- and zooplankton were simultaneously assessed at a large spatial scale and compared with respect to environmental variables and anthropogenic inputs. Factor analysis revealed the effects of load of inland waters, seasonality, water turbulence and hydrocarbon pollution on the planktonic components and zooplankton variability in port sectors characterized by different depths and uses.

REVIEWER #1: MPB-D-19-01921

R#1.1: This manuscript presents the results of nine plankton surveys conducted at three Mediterranean ports, during February, May and September 2012, in order to evaluate the effects of physical and chemical water variables on bacterioplankton, phytoplankton and zooplankton. Literature information on the above plankton components combined, in ports, is missing. There are not innovative approaches, but the work is well designed and has been carried out meticulously.

ANSWER: Literature including all the three components in harbours on the Mediterranean Basin scale is absent. Some references were added (Pestoric et al., 2018; Bernàt et al., 1994), which analysed the three components of plankton.

R#1.2: The manuscript is long and sometimes heavy to read in some parts (results, discussion) due to various unnecessary details especially in the discussion. However, it is well organized and written. My suggestion is the shortening/revision of discussion having in mind that port environment is very unstable and fluctuating, and cannot be directly compared with most other ecosystems. Also, this environment is especially vulnerable to meteorological conditions.

ANSWER: We agree with these considerations. In the discussion, we removed the open sea as comparison and revised the other comparisons. Moreover we shortened some sections in paragraph 4.1 and strongly reduced paragraph 4.2 (former page 10, lines 5-7; former page 11, lines 14-25 & 34-35).

R#1.3: Another problem is the formulation lack of a clear synthetic conclusion on the main and more innovative results, in other words, the novel contribution of this work with respect to previous similar ones.

ANSWER: We thank the referee for the constructive comments. Conclusions were re-elaborated, shortened and better focused. References were removed.

R#1.4: In general, English need improvement, with special attention in the discussion section.

ANSWER: English was carefully checked and revised, particularly in the discussion section.

R#1.5: P 7, L 51-52 Remove this sentence, unnecessary.

ANSWER: former Page 7, Lines 51-52: The text was removed according to reviewer suggestion.

R#1.6: P 11, L 7-8 Replace this sentence, for example "It might be related with the fact that Cagliari port receives freshwater inputs and/or with the advection from neighbouring areas".

ANSWER: former Page 11, Lines 7-8 The text was revised according to reviewer suggestion.

R#1.7: P 11, L 35-36 No meaning to compare offshore waters with port waters.

ANSWER: the whole paragraph 4.2 was modified accordingly and shortened also according with point #1.2.

R#1.8: P 11, L 45 Have always in mind that plankton advection is very important for the stations close to the port entrance.

ANSWER: former Page 11, Line 45 The text was integrated according to reviewer suggestion and a reference was added.

R#1.9: P 12, L 4-5 you cannot comment the offshore currents of course they do not affect any of the ports. Just, simply state that the variable plankton advection cannot be measured.

ANSWER: former Page 11, Lines 2-6: The sentence was removed so that the necessity of considering here the plankton advection has come to nothing. Anyway the possible effect of this variable was considered throughout the text following the Reviewer suggestion.

R#1.10: In general, throughout "discussion", revise the various comparisons.

ANSWER: The text was revised according to reviewer suggestion.

R#1.11: P14 L 49 - This part needs revision, it should be more concise. It should report original synthetic considerations emerged from the Discussion and highlight the most original findings of the work. Exclude repetitions of results which have already been discussed. Remove references in the Conclusions.

ANSWER: We perfectly agree with the referee. The text was corrected also according to points R#1.3 and R#2.22.

R#1.12: Figure 1 - Better to follow the same order as in results (always the same throughout text); Cagliari, El Kantaoui, Heraklion "Maps of the three Mediterranean ports and sampling stations. a) Port of Cagliari (Italia, Sardinia) and inlets of freshwaters (arrows); b) Systems of lagoons and canals surrounding the Port of Cagliari (in the box); c) Port of El Kantaoui (Sousse, Tunisia) d) Port of Heraklion (Crete, Heraklion)."

ANSWER: The figure 1 and its figure legend were revised according to the reviewer suggestion.

REVIEWER #2: MANUSCRIPT NUMBER: MPB-D-19-01921

The MS addresses the comparison of three Mediterranean ports in relation to the plankton and the variables that affect it. In my opinion, it's an interesting topic; the use of (zoo)plankton as an indicator of water quality in closed water masses and small systems, which, sadly, in most of the cases are ostracized. The paper is not too long and adequately understandable. However, the MS can be improved. Globally, authors have attempted to

i) assess zooplankton abundance and composition variability,

ii) to define its relationships with the other planktonic components (bacterioplankton and phytoplankton) and with the water physical and chemical parameters, and

iii) to identify the potential impacts of coastal pressures, port activities and linked infrastructures on the planktonic network.

R#2.1: In my opinion, in the case of the objectives i) and ii) they are not plenty fulfilling. Zooplankton composition and abundance are not shown in totality (Table S1), and the showed results/data and the comments in results/discussion mismatch (for example copepod taxa). There is no problem if grouping the taxa, but the same groups should be used in the analyses (Table 3) and graphs.

ANSWER: We thank the referee for this constructive comment. Tables 2 and 1S were revised and homogenised to be more informative and respondent to the data reported in the text. To do this we divided Table 2 in Table 2a (physical and chemical variables) and Table 2b (biological variables). Table 1S was divided in Tables 1S (physical and chemical variables, bacterio- and phytoplankton) and 2S (all the replicates of zooplankton). All the inconsistencies were corrected. Caption of Figure 2 was modified accordingly. Considering statistical analysis, our choice of performing the analysis only with taxa that reached a certain abundance ($N \geq 50$) was dictated by the fact that introducing in the Factor Analysis the species that occurred in too few numbers and samples increases noise without adding information. That is a common practice when applying Factor Analysis.

R#2.2: Regarding objective ii), the relationships between zooplankton and the other planktonic components (bacteria & phyto) are scarcely analysed, the majority of arguments are from literature but, in my view, the authors don't make the most of the data to establish these relationships/influences.

ANSWER: We agree with the referee. The relationship between zooplankton and the other planktonic components was better focused in the text. More specifically, sentences were added at the end of paragraphs 4.1 and 4.2 of Discussion (former page 11 line 9 and page 12 line 43) to highlight these relationships. This was further considered in Conclusions that were modified accordingly.

R#2.3: And regarding this, why AF analysis? what are the reasons/advantages of choosing AF and no other multifactorial analysis? Highlight this point, please.

ANSWER: We agree with the reviewer on this point. There are many possible ways to analyse a dataset. In this study, we selected factor analysis based on several advantages over other multifactorial analysis. An important advantage of AF is that it can be used as exploratory technique to search for patterns among a set of variables (our objective ii: to define relationships among planktonic components and with the water physical and chemical parameters), and as such it does not require to pre-define which variables affect which variables and how (we are exploring, we are searching that knowledge to define the relationships). Another important advantage of AF is that it does not require (multi)normality because no hypothesis will be tested (we are exploring, we still don't have a hypothesis to test). The exploration would eventually lead to hypothesis that then may be tested, and the need for this is due to the lack of previous studies on the subject in ports that consider many variables simultaneously.

Page 6 lines 23-24: The text was modified to synthetically clarify the concept.

R#2.4: 2.3.4 Zooplankton. Page 5 Line 60: "... five vertical tows were performed..." and Page 6 Line 3: "The individuals in the five replicates...". I don't understand well if the five replicates were analysed separately (and in this case, then the mean was calculated) or all of them were grouped in one lonely sample and then it was analysed. Please, clarify this point.

ANSWER: The text was revised according to reviewer suggestion in the Material & Methods in the sections 2.3.4 "Zooplankton" and 2.4 "Statistical analysis" to better clarify this point. Each replicate was sorted and analysed separately. The data of each replicate are now reported in Table 2S (Supplementary material), while in Table 2 the means were reported to simplify the readability of the results.

R#2.5: 2.3.4 Zooplankton. Page 6 Line 4-5: "... The main taxa were identified to the lowest possible taxonomic level, ...". However, in the results, only the main groups are shown, and in the case of the copepods at genus level.

ANSWER: The sentence was removed and the text was revised according to reviewer suggestion. The main groups were reported in Table 2 and Table 2S.

R#2.6: 2.4. Statistical analysis. Page 6 Line 21. "The relationships among factor analysis.... to identify the... variability of the data..." What is the reason to perform factor analysis? Why no RDA, for example, to relate environmental and biotic variables?

ANSWER: Factor analysis is an appropriate technique to simultaneously explore relationships among variables that are simultaneously interacting, without assuming pre-defined cause-effect relationships. The AF permits to highlight the multiple relationships and weight their importance in the system.

In this study, we considered the planktonic environment as a whole, including organisms and water physical-chemical variables. The AF allows including both, organisms and environmental variables, which in the reality do interact. Consider that not only the physical chemical variables influence organisms, but that organisms influence the water characteristics (e.g., phytoplankton influences chlorophyll content, water transparency, ...). The AF helps to focus on the interpretation of the relationships.

On the other hand, Redundancy Analysis (RDA) requires to pre-define which variables affect which variables, hence we considered it not appropriate for our objectives. RDA is similar to canonical correlation analysis but allows the user to derive a specified number of synthetic variables from one set of (independent) variables that explain as much variance as possible in another (independent) set. It is a multivariate analogue of regression. As such it requires a pre-definition of dependent and independent variables, and also multinormal distribution of the variables.

Former Page 6 lines 23-24 The text was modified to synthetically clarify the concept (as point R#2.3).

R#2.7: Which is/are the latent variable(s) (unobserved variables) associated with the FA?

ANSWER: The unobserved/latent variables are the interpretation of the four factors extracted, as indicated in the last line of Table 4 and explicitly addressed in the discussion section: Factor1 is freshwater input and nutrient loading, Factor2 is zooplankton variability, Factor3 is seasonality and water turbulence, Factor4 is anthropogenic activities.

R#2.8: 3.1. Description of the physical-.... Page 6 Lines 49-50: "The only exception was the concentration of AHs in The Kantaoui marina...." And Nitrate and Silicate in Heraklion? Is the same result.

ANSWER: We agree with the referee. We corrected the text, former Page 6, Lines 49-50 from the original "The only exception was the concentration of AHs in the El Kantaoui marina, which was not significantly different from the other two ports." in "The only exceptions were the concentrations of AHs in El Kantaoui and nitrate and silicate in Heraklion (Table 3)."

R#2.9: 3.1. Description of the physical-.... Page 7 Lines 13-15: "...The overall variance accounted by this model was only 18% of the total variance...." Please, join some way with the previous sentence. It's not clear the reference to only PAHs.

ANSWER: We agree with the referee. We correct the text, former Page 7, Line 13-15: "The overall variance accounted by this model was only 18% of the total variance, as shown by the low coefficient of determination ($r^2=0.18$, Table 3)." in "The overall variance accounted for by PAHs model was 18% of the total variance, as shown by the lower coefficient of determination of PAHs ($r^2=0.18$, Table 3) as compared to the other variables ($r^2>0.70$, Table 3)."

R#2.10: 3.2. Planktonic biota in the three ports. Page 7 Line 27: (ANOVA, Table 3). Please, delete ANOVA

ANSWER: former Page 7, Line 27: "ANOVA" was removed in the text

R#2.11: 3.2. Planktonic biota in the three ports. Page 7 Lines 51-55: I have not clear where do the authors obtain these density values: 36,659, 5,791, 63,228, 2,022, 1,652 and 1,175 ind. m⁻³. Not in table 2 nor Table 1S appear.

ANSWER: Former Page 7, Lines 51-55. We agree with the referee, reported data are inconsistent with Tables since they derived from not shown raw data. Accordingly, and also following the request of Reviewer 1, the text at lines 51-52 was removed and lines 51-55 corrected accordingly with Tables 2 and 2S.

R#2.12: 3.2. Planktonic biota in the three ports. Pages 7 & 8 Lines 60 & 1-2: "Overall, over 70 zooplankton taxa were identified ... (Table 1S, Supplementary Material)." the taxa (70) are not shown, only the main zooplankton groups.

ANSWER: We agree with the referee. The inconsistency was due to the fact that many taxa have very low abundance and were grouped. The text was revised in order to clarify this point avoiding discrepancies. Former Page 7 & 8, Lines 60 & 1-2 were modified subsequently: "Overall, 39 zooplankton groups were recognized, (Table 2b)."

R#2.13: 3.2. Planktonic biota in the three ports. Pages 8: Figures 2a & 2b and related text. I don't understand the difference between density values, there is a mismatch between tables (2 and S1) and figure 2. For example, in Fig 2a in H3 on February for total zooplankton almost 4000 ind m⁻³ are recorded, however, in table S1 supplementary material, for this station and month (1H3) the value is 2020. And similar for the rest.

ANSWER: We agree with the referee. There was an error in the data representation and data reported. Figure 2a was corrected

R#2.14: 3.2. Planktonic biota in the three ports. Pages 8 Lines 37-41: "The seasonal variation of single.... (Table 3)." The own variability of the zooplankton taxa/groups (e.g. seasonal cycles and distribution) can imply equality if total zooplankton is analysed. Hence, if the goal is to know the effects (of different variables) on the biota, the analysis on the different taxa, or at least, the main groups (as the phytoplankton, for example) is necessary. In this sense, the factor 2 (Factor analysis) account for the different groups of zooplankton, so could be interesting to analyse possible differences.

ANSWER: The sentence was modified. Factor2 identified the zooplankton groups with similar patterns of variability, hence the analysis of Factor2 is in itself the analysis of those zooplanktonic groups (of the different zooplanktonic taxa grouped by common patterns). This point was better clarified in the text.

R#2.15: 3.3. Linking abiotic condition and planktonic... Table 4 & Figure 3. The results of table 4 are from the analysis of data as a whole, while in Figure 3 are shown by space (port and stations within) and by time. However, this distinction and the relationship between both results is not specified in Material and Methods.

ANSWER: Table 4 and Figure 3 were built from the same set of data as reported at the beginning of paragraph 2.4 of M&M. We better clarify this point at the end of paragraph 2.4: "The values of the factors calculated for each station and sampling period in the three investigated ports were graphically analysed."

R#2.16: 4.1. Differences in freshwater inputs.... No differences on zooplankton between ports have been recorded but there are on environmental variables and bacteria and phytoplankton. These differences are related to freshwater inputs as consequences of the different hydrography of the ports, mainly Cagliari. In this sense, and taking account that samples were obtained in surface, what is the structure of the water column in the ports? Stratified system? Mixed water column? Because at a certain depth (zooplankton habitat) the variables can become similar and this one explains some results. Have the authors studied the water profiles?

ANSWER: We better focused in the discussion section the differences among the studied ports in the zooplankton component. More specifically, Factor1 highlighted marked differences in the zooplankton (Cladocera for example) distinguishing Cagliari port from the other two ports. Moreover, differences among ports in the taxa included in Factor2 was also found and the text was better clarified in the Discussion section.

We did not study the water profiles, as the depths at the sampling stations and in the whole ports were overall small, particularly in the stations for docking of leisure boats in all ports (stations 1) and specifically in El Kantaoui Port. Moreover, the water in some areas was frequently mixed by boats and vessels passages.

R#2.17: 4.2 Consumer dynamics. What do the authors mean by "consumer"? All zooplankton taxa are consumers. And on the other hand, sampling periodicity (three months of one year) is not enough to study the dynamic.

ANSWER: We agree with the referee. We corrected "Consumer dynamics" in "Zooplankton variability" all over the text.

R#2.18: 4.2. Consumer dynamics. Page 11 Lines 33-35: "The total number of individuals observed in literature open sea gradient". What literature?, Previously cited?. Besides, in the previous paragraph, the authors reported overall densities at different Mediterranean sites, that in my opinion, they are not very relevant for the discussion (can be summarized), but in this sentence, no results are shown, which could be more interesting for the comparison (similarities) among the sites and to explain the reasons.

ANSWER: We thank the referee for the constructive comment. We revised the text in 4.2 Section, that was simplified also according to the suggestions of Referee 1. The whole data on the open sea gradient reported from literature were removed.

R#2.19: 4.2. Consumer dynamics. Page 11 Lines 51-60 & Page 12 Lines 1-19: The paragraph deals with the abundance differences between ports/stations. However, the only statistical comparison for total zooplankton of Cagliari stations (Table 3b) is not significant (for taxa and other stations are not shown).

ANSWER: The text refers to the abundance of the specific zooplankton groups included in Factor 2 of Factor Analysis (as reported in Table 4 and shown in Figure 3), not to the total abundance of zooplankton (as shown in Table 3). We clarified the statements accordingly, changing the original "the majority of zooplankton taxa (Factor2, Table 4)" in: "specific groups of zooplankton (as shown by Factor2, Table 4)"

R#2.20: 4.3. Seasonality and water turbulence. Page 13 Lines 14-18: "Concerninggenus Oithona ... They have ... negatively influenced by water temperature (Wang et al., 2017)". Oithona is a broad genus with numerous species that present divergent relationships with temperature and in the case of the paper of Wang et al., they only sampled in summer and the negative influence of the temperature was, mainly, on O. plumifera.

ANSWER: The sentence was corrected and the concept clarified with more precise references to cited papers.

R#2.21: 4.4. Impacts of anthropogenic activities. Page 14 Line 32 "... (Pearson $r_2 = 0.34$; $r_2 = 0.44$) ... ". Please indicate also p-values

ANSWER: The p-values were indicated in the text according to the reviewer suggestion

R#2.22: Conclusions. Maybe, it's a journal/editor decision but, in my opinion, the conclusions should avoid citations. Conclusions are a summary of findings from the study and from literature. The comparison, justifying, interpretation of the findings with the bibliography should have been completed in the discussion.

ANSWER: We thank the referee for the constructive comments. Conclusions were re-elaborated, shortened and better focused also according to suggestions of the Referee 1.

R#2.23: Table 1. Unnecessary, it is explained in the text, and the water depth data can also be added on the text.

ANSWER: We added this Table and consider it useful since together with Figure 1 it makes easier for the reader to follow the organization of the sampling areas, localize the single stations through geographical coordinates (that is difficult to insert in the text) and define their uses. Moreover, explanations of labels of Figure 2 and 3 have a reference to Table 1.

R#2.24: Table 2: Please indicate Proportion of Holoplankton

ANSWER: We added the proportion of Holoplankton

R#2.25: Table 3. 1-way Anova table 3: In my opinion, this (part of) table is irrelevant, the information can be added in the text as you do with Heraklion. Anyway, what means A or B?

ANSWER: We corrected "A" and "B" in "a" and "b". We considered this part of the table relevant since it shows the variables statistically different among stations within the Cagliari port. This allows the identification of the significant abiotic variables and the different inputs in this complex port. We think therefore that the table should be presented with the 1-way-ANOVA. We integrated the text to better present data shown in this part of the Table, also according to point #2.9.

R#2.26: REFERENCES

R#2: Page 10, line 58: Heneash et al., 2014 is not in References.

R#2: Page 17, lines 54-58: "Heneash, A.M.M., Tadrose, H.R.Z., Hussein, M.M.A., Hamdona, S.K., Abdel-Aziz, N., Gharib, S.M., 2015. Potential effects of abiotic factors on the abundance and distribution of the plankton in the Western Harbour, south-eastern Mediterranean Sea, Egypt. Oceanologia 57(1), 61-70. <https://doi.org/10.1016/j.oceano.2014.09.003>" is not in the text
References. Page 20, lines 9-10: "Roberts, D.A., 2012. Causes and ecological effects of resuspended contaminated sediments (RCS) in marine environments. Environment International, 40(1), 230-243. <https://doi.org/10.1016/j.envint.2011.11.013>" is not in the text

ANSWER: Page 10, Line 58 The citation in the text was corrected in "Heneash et al., 2015"

R#2: Page 14, lines 4 & 7: Bullita, 2016 is not in References.

R#2: Page 16, lines 18-22: "Bullita, E., Pani, A., Tamburini, E., 2016. Dottorato di ricerca in Sviluppo e Sperimentazione di Farmaci antinfettivi Ciclo XXXVIII Characterization and comparison of microbial communities

from different tourist ports in Mediterranean Sea and evaluation of applicability of bioremediation treatment. PhD. <https://doi.org/10.1016/j.bmc.2015.08.028>" is not in the text References.

ANSWER: Page 14, Lines 4 & 7: The citation in the text was corrected in "Bullita et al., 2016"

R#2: Page 14, lines 11-12: Roberts et al., 2012 is not in References.

R#2: Page 20, lines 9-10: "Roberts, D.A., 2012. Causes and ecological effects of resuspended contaminated sediments (RCS) in marine environments. Environment International, 40(1), 230-243. <https://doi.org/10.1016/j.envint.2011.11.013>" is not in the text

ANSWER: Page 14, Lines 11-12: The citation in the text was corrected in "Roberts, 2012"

R#2: Page 14, line 26: Duran et al., 2016 is not in References.

R#2: Page 17, lines 30-32: "Duran, R., and Cravo-Laureau, C., 2016. Role of environmental factors and microorganisms in determining the fate of polycyclic aromatic hydrocarbons in the marine environment. FEMS Microbiol. Rev. 40, 814-830. <https://doi.org/10.1093/femsre/fuw031>" is not in the text References.

ANSWER: Page 14, Lines 30-32: The citation in the text was corrected in "Duran and Cravo-Laureau, 2016"

*Highlights

- Ports are impacted coastal areas and their waters need to be monitored
- Plankton communities are abundant and composite in port waters
- Bacterioplankton, phytoplankton and zooplankton constitute a complex network
- Plankton networks depend on nutrient load, water turbulence, seasonality and hydrocarbons
- Plankton network usefully contribute to the monitoring of port areas

VARIABLES AFFECTING THE PLANKTON NETWORK IN MEDITERRANEAN PORTS

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Abstract

1 Attention on port waters is increasing since these economically important infrastructures are embedded in
2 the coastal environment and their management needs to be considered in the monitoring programmes of
3 coastal ecosystems. To implement the sustainable development (blue growth) of port areas, a general
4 knowledge on the ongoing processes in their waters needs to be obtained, considering both abiotic and
5 biotic variables. The present study aimed at inspecting the relationships among plankton components to
6 provide insights into ecology of ports. Seasonal samplings were carried out in three Mediterranean touristic
7 ports where bacterio-, phyto- and zooplankton were simultaneously assessed at a large spatial scale and
8 compared with respect to environmental variables and anthropogenic inputs. Factor analysis revealed the
9 effects of load of inland waters, seasonality, water turbulence and hydrocarbon pollution on the planktonic
10 components and zooplankton variability in port sectors characterized by different depths and uses.
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14 **Keywords:** plankton, zooplankton community, Mediterranean Sea, port, anthropogenic impact,
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1. Introduction

1 In recent decades international policies have converged toward a coherent management of coastal areas,
2 integrating land and sea ecosystems in the coastal zone and their interactions (Newton and Icelly, 2008) on
3 one side, and environmental and economic issues on the other. The key concept “sustainability” represents
4 the main focus of many of the current policies, but a general failure is observed, primarily due to a lack of
5 coordination among the different stakeholders (McGuire and Perivier, 2011) and the conflict between
6 economic development and environmental quality. The European Community established a framework of
7 actions and provided directives to define and improve the ecological water quality (Water Framework
8 Directive, WFD 2000/60/EC; Marine Strategy Framework Directive, MSFD 2008/56/EC; Maritime Spatial
9 Planning Directive, MSPD 2014/89/EU) with the aim of protecting the marine ecosystems and ensuring the
10 delivery of their important ecological services.

11 In the restricted space of the Mediterranean coasts, the number of ports and marinas is constantly
12 increasing, competing with bathing waters and protected areas (Piante and Ody, 2015). Port waters are not
13 only directly linked to marine waters through the port entrance, but also to inland waters through inlets
14 and discharge pipes. Therefore, the expansion of port areas could exponentially contribute to the decline of
15 coastal water quality (Durrieu de Madron et al., 2011), potentially affecting the ecosystem services
16 provided (e.g. fisheries, biodiversity, water regulation, cultural benefits). Despite their strategic role in
17 coastal development, the main controls carried out in ports and adjacent areas are those related to
18 chemical emissions and consequent pollution in water and sediments (Peris-Mora et al., 2005; Di Vaio and
19 Varriale, 2018) as well as sanitary aspects linked to microbial pathogens (EEA, 2017), as basic controls to
20 protect human health. Nevertheless, the most suitable Biological Quality Elements (BQEs:
21 macroinvertebrates, angiosperms, macroalgae, phytoplankton and fish) as defined by WFD (2000/60/EU)
22 for the environmental impact assessment in natural water bodies have also been suggested for ports, but
23 not yet extensively implemented (Hering et al., 2010; Ondiviela et al., 2013).

24 In ecological monitoring, the analysis of sediments and associated benthic biota are currently the most
25 used approaches to develop ecological quality indexes (Quintino et al., 2006; Teixeira et al., 2016). The
26 stable and long-life characteristics of benthic communities, whose structure accumulates with time
27 nutrients and contaminant loadings in the sediments, strikingly differ from the greater variability of the
28 planktonic biota (Caroppo et al., 2013). Under the EU directive framework, water quality has been
29 therefore analysed using indices based on benthos (Diaz et al., 2004) and standards have been developed
30 for coastal waters (WFD, 2000/60/EU). Only chlorophyll-*a* has been extensively used as the most shared
31 ecological quality indicator of the water column (Dimitriou et al., 2015).

32 From an ecological point of view, bacterioplankton, phytoplankton and zooplankton are the main
33 planktonic components present in the water column in terms of abundance, biomass, diversity, trophic
34 networks and ecosystem services provided (Beaugrand et al., 2010; Siokou-Frangou et al., 2010; Tweddle et
35 al., 2018). Planktonic taxa show a large spectrum of size, trophic and ecological roles and have been rarely
36 considered together as descriptors of the marine ecological quality (Caroppo et al., 2013). In the water
37 column, physical-chemical and biotic parameters undergo higher variability at shorter temporal scales than
38 benthic ones, with regard to seasonal changes and inflow/outflow from the connected marine and inland
39 water bodies. The planktonic taxa can thus be perceived as a “moving interface”, rapidly reacting to the
40 environmental variations and connected with the more stable benthic communities. Thanks to their
41 biological properties, the planktonic communities have been proposed as early warning indicator of several
42 types of impacts. More specifically, bacterioplankton abundance and activities have been acknowledged as
43 sensitive sentinels of environmental changes (Munawar and Weisse, 1989; Caruso et al., 2016a; Caruso et
44 al., 2016b), phytoplankton blooms have been related to eutrophication processes (Karydis, 2009) and
45 several other pressures (MSFD criterion 4.1.), while zooplankton variation has been linked to regime shifts
46 in ecosystem state (Pace et al., 2013), climatic changes (Roemmich and McGowan, 1995; Beaugrand, 2005;
47 Sundstrom et al., 2017) and pollution (Uriarte et al., 2016). In this perspective, an all-inclusive control
48 strategy is desirable in order to monitor the ecological status of water bodies (WFD, 2000/60/EC).
49 Considering the continuity of coastal areas (which include marine protected areas, coastal lagoons, river
50 mouths, bathing areas, tourist and commercial infrastructures, marinas and harbours, etc.), an in-depth
51 knowledge of the planktonic components is an essential step to assess the mutual influence between linked

1 water bodies, which are potential sources of abiotic and biotic variability (e.g. eutrophication,
2 hydrocarbons, heavy metals, presence of euryhaline species, etc.).

3 Based on these concepts, a pilot study using an integrated planktonic approach was needed to build a
4 knowledge framework on trophic processes of the networks prevailing in port waters, since morphology,
5 hydrodynamics and human activities in harbours (heavily modified water bodies, WFD, 2000/60/EC) may
6 specifically affect bacterio-, phyto-, and zooplankton as well as their relationships (Caroppo et al., 2013).

7 The present study was conducted under the framework of the ENPI-CBC MED project MAPMED
8 (Management of Port areas in the Mediterranean Sea Basin), a multidisciplinary project aimed at improving
9 the environmental sustainability in Mediterranean tourist ports (Zakhama-Sraieb et al., 2016;
10 Chatzinikolaou et al., 2018; Massi et al., 2019; Vitali et al., 2019). In this study, the planktonic components
11 of three Mediterranean tourist ports in different periods of the year, related to the tourist season, were
12 analysed with the following specific aims: i) to assess zooplankton abundance and composition variability,
13 ii) to define its relationships with the other planktonic components (bacterioplankton and phytoplankton)
14 and with the water physical and chemical parameters, and iii) to identify the potential impacts of coastal
15 pressures, port activities and linked infrastructures on the planktonic network. The different parameters
16 (physical, chemical and biological) were analysed by means of a multivariate analysis in order to compare
17 and interpret the ecological relationships in the water column of the investigated ports.

20 **2. Material and Methods**

21 **2.1. Brief description of the three ports**

22 Among the three studied ports (Figure 1), Cagliari (Sardinia, Italy) and Heraklion (Crete, Greece) are both
23 commercial and touristic harbours with a larger surface (2.07 km² and 0.87 km² respectively) than the
24 artificial marina (0.04 km²) of El Kantaoui (Sousse, Tunisia), where activities linked to tourism and fishing
25 are operated. The three ports have an inner and more protected shallow area hosting small leisure and/or
26 fishing boats; from there to the port entrance, higher depths and larger infrastructures (quays or docks)
27 allow for activities involving bigger ships and cargoes in Cagliari and Heraklion.

28 Within each port, three to five sampling stations were selected to achieve a good spatial coverage of the
29 whole port area and represent discrete sectors dominated by specific port activities, potentially exposed to
30 different impacts (Table 1, Figure 1). In detail, the Cagliari port hosts quays for the anchoring of small
31 leisure boats (station C1), military navy, commercial and passenger ships (stations C2, C3, C4, respectively).
32 The port entrance (station C5) faces south-west. The opening of Molentargius Lagoon loads brackish waters
33 into the port main basin at station C1, while a canal running along the east side of the Santa Gilla Lagoon
34 drains wastewater of urban runoff in the proximity of station C4 (Figure 1a and b). The marina of El
35 Kantaoui is the second tourist port in Tunisia and hosts small and medium-size leisure boats (station E1).
36 Only a small part of the marina is dedicated to other activities, as shipyard and fuel supplying (station E2).
37 The port entrance (station E3) faces south-east (Figure 1c). The Heraklion port is constituted by the Old
38 Venetian Harbour, hosting leisure and fishing boats (station H1), and the new Ferry Port, where passenger
39 and cargo ships (stations H2, H4, respectively) are hosted along with a shipyard (station H5). Heraklion port
40 entrance (station H7) faces east (Figure 1d). Discharges of sewage effluents from urban activities have been
41 documented in the three port areas (Chatzinikolaou et al., 2018; Massi et al., 2019; Vitali et al., 2019).

42 **2.2. Sampling campaigns**

43 Three sampling campaigns were performed in 2012 during winter (February), spring-early summer at the
44 beginning of the tourist season (May), and late summer-autumn at the end of the tourist season
45 (September). The samplings were performed during the day, from 9 am to 6 pm, to avoid the effects
46 deriving from day-night plankton migrations. This sampling design does not account for the planktonic taxa
47 life cycle variation (from few days, to months and years, depending on the taxon), but was conceived to
48 capture the major environmental variations due to both climatic factors and human activities and the
49 consequences they may have on the planktonic communities. In the Mediterranean region, winter and
50 spring are normally rainy seasons, with lower temperatures and higher fresh water load, which accounts for
51 a slightly lower salinity along the coasts (Mehta and Yang, 2008); summer is warm and dry, with major
52 effects on water salinity in early autumn because of the increased evaporation. The tourist season starts in

1 late spring and ends in early autumn, resulting in water scarcity, intense use of coastal facilities, high
2 volume of marine traffic, as well as increased discharge/spill of wastewaters and pollutants (e.g. sewage
3 effluents, lubricating oils, fuel oils and combustion products).

4 **2.3. Sampling procedures and laboratory analyses**

5 *2.3.1. Physical and chemical variables*

6
7 The physical properties of surface water (temperature, salinity, oxygen and pH) were measured on board
8 using a 3420 WTW multi-meter. Three replicate samples (5 L each) of surface seawater were collected at
9 each station during each sampling campaign and used for chemical analyses (Table 1S - Supplementary
10 material).

11 For the chemical analyses of inorganic nutrients (NO_2 , NO_3 , NH_4 , PO_4 , SiO_2), particulate organic carbon
12 (POC) and chlorophyll-*a*, the seawater samples were filtered immediately after collection through
13 Whatman GF/F filters (47 mm). Filters were stored at -20°C and used for the determination of chlorophyll-*a*
14 according to the fluorometric method of Yentsch and Menzel (1963) and Arar and Collins (1992). The
15 filtered water samples were stored at -20°C and used for the determination of NO_2 , NO_3 , PO_4 , SiO_2 ,
16 following the techniques proposed by Strickland and Parsons (1972) and for NH_4 , those by Ivančić and
17 Degobbi (1984).

18 For hydrocarbons, samples of 1 L of unfiltered water were extracted with hexane spiked with surrogate
19 standards of aliphatic hydrocarbons (AHs) and polycyclic aromatic hydrocarbons (PAHs). The extracts were
20 concentrated, fractionated and the concentrations of AHs and PAHs were measured using gas
21 chromatography-mass spectrometer (Agilent 6890 gas chromatograph interfaced with mass spectrometer).
22 Determination of heavy metals in water samples was performed by Inductively Coupled Plasma
23 Spectrometry (ICP-OES, Perkin Elmer Optima DV 7000). Concentrations resulted below the detection limits
24 of the analytical method in all samples for As, Cd, Cr, Ni, Pb, Sb, and V, and in more than 50% of the
25 samples for Cu; therefore metals were not further included in the analyses (MAPMED Consortium, 2013).

26 *2.3.2. Bacterioplankton*

27 Water samples were collected at the surface (within 1 m depth) using sterilized 15 L low-density
28 polyethylene collapsible carboys (washed with 10% bleach and rinsed with sterile MilliQ water). Samples
29 were immediately fixed with filtered formaldehyde (final concentration 1.8%) for 1 h at 4°C . One mL
30 aliquots were filtered onto black polycarbonate membranes (0.2 μm pore size, 25 mm diameter), rinsed
31 with ultra-pure water, air-dried, and transported to the laboratory at room temperature (Beardsley et al.,
32 2008). The filters were stained with DAPI (10 $\mu\text{g mL}^{-1}$) for 5 minutes, then washed six times with ultra-pure
33 water and six times with ethanol 80%. Filters were mounted onto microscope slides using UV-transparent
34 fluorescence-free immersion oil. Cells (diameter $< 20 \mu\text{m}$) were counted via epifluorescence microscopy
35 (Olympus BX51) equipped with a mercury burner power supply unit (OLYMPUS U-RFL-T) in five fields per
36 filter on three replicate filters for each station and each sampling period.

37 *2.3.3. Phytoplankton*

38 Samples of 250 mL of water were collected at the surface (within 1 m depth) at each station and each
39 sampling period, fixed with neutralized formalin (final concentration 1%) and stored in dark glass bottles.
40 Subsamples of variable volumes were observed under an invertoscope (Zeiss IM35, ph. c., 40x) after
41 sedimentation, following standard methods (Zingone et al., 2010).

42 Phytoplankton taxa were identified and assigned to the classes of diatoms, dinoflagellates and
43 coccolithophores; cryptophytes, chlorophytes, cyanobacteria and nanoflagellates, which could not be
44 identified further, were included in the mixed group labelled as "other phytoplankton".

45 *2.3.4. Zooplankton*

46 An Apstein net for zooplankton (200 μm mesh width, 40 cm mouth diameter, 1 m net length) was used and
47 five vertical tows were performed (Zunini Sertorio, 1990; Camatti and Ferrari, 2010) at each station and
48 during each sampling period in the three ports, avoiding the net touching the bottom. The volume of
49 filtered water was calculated as: $V = \text{mouth surface} \times \text{station depth}$ and was used to estimate the densities
50

of zooplankton (ind m⁻³). The samples were stored in 8% normalized formalin and analysed. The individuals were sorted, counted and identified under a stereomicroscope using a Bogorov counting chamber for zooplankton (40 mL). The counts and taxonomic identifications of the five independent zooplankton replicates were retained as separate samples for statistical analysis (Table 2S - Supplementary material).

2.4. Statistical analysis

For each station and each sampling period in the three ports, a data matrix was constructed including the values of physical-chemical parameters, total bacterial counts, densities of total phytoplankton and its groups (diatoms, dinoflagellates, coccolithophores, "other phytoplankton"), and counts of total zooplankton and its groups (five replicates, Table 1S). The bacterioplankton, phytoplankton and zooplankton counts were log transformed to approach normal distribution. The same set of physical-chemical, bacteria and phytoplankton data collected in a station was used for the five zooplankton replicates collected in the same station.

For univariate analysis, 2-way-ANOVAs with port and sampling period as main effects and 1-way-ANOVAs by stations within each port were performed (Statistical Analysis System, SAS, package version 9.4). When ANOVA detected significant effects ($p < 0.05$), the Scheffé multicomparison of means was performed. The Scheffé test was selected because it can compare groups with different number of observations (Scheffé, 1959).

The relationships among variables were evaluated with the multivariate technique of factor analysis (SAS, package version 9.4) to identify the environmental variables and groups of planktonic taxa accounting for the main variability of the data without assuming pre-defined cause-effect relationships (Kim and Mueller, 1978; Milstein, 1993; Nourisson et al., 2018). In this analysis we included the physical-chemical variables, total bacterial counts, densities of the four phytoplankton groups and densities of the zooplankton taxa that presented at least 50 ind m⁻³ in the overall database. Among the several available techniques to extract factors, the Principal Component Analysis (PCA) calculated from the correlation matrix among variables was selected, which allows the factor analysis to be applied as exploratory tool, without requiring a normal distribution of all the variables included in the data matrix (Kim and Mueller, 1978). The method computes the linear combination of the original variables, which accounts for as much of the variation contained in the samples as possible, called first factor (Factor1). The second factor (Factor2) is the second linear function of the original variables, which accounts for most of the remaining variability, and so on. The factors are independent one from another, have no units and are standardized variables (normal distribution, mean=0, variance=1). The value of a factor in a given sample is the result of the sum of all the variables included in the factor calculation, each one multiplied by a coefficient. The values of the factors calculated for each station and sampling period in the three investigated ports were graphically analyzed. The coefficients of the linear functions defining the factors were used to interpret their meaning, considering the sign (+/-) and relative size of the coefficients as an indication of the weight of each variable.

3. Results

3.1. Description of the physical-chemical variables in the three ports

Temperature and salinity were significantly lower and the majority of the other variables were significantly higher in Cagliari port than in El Kantaoui and Heraklion ports (Table 2a, Table 3 for ANOVA). The only exceptions were the concentrations of AHs in El Kantaoui and nitrate and silicate in Heraklion (Table 3). No significant difference was found in pH among ports. As expected in the Mediterranean region, temperature increased from winter (February) to early summer (May) to late summer (September) and dissolved oxygen (DO) was significantly lower in September than in February and May. The variable pH was significantly higher in September than in February, with intermediate values in May. Both POC and chlorophyll-*a* were significantly higher in May than in February, with intermediate values in September (Table 2a, Table 3 for ANOVA). On the overall dataset, polycyclic aromatic hydrocarbons (PAHs) were significantly higher in February than in the remaining months. Overall, the variability among ports was higher than the variability among sampling periods, since 12 of the 13 analysed physical-chemical variables presented significant differences among ports and only 6 among periods (2-way-ANOVAs, Table 3).

1 The variability among stations within each port was lower than the variability among ports and among
2 periods, as indicated by the low number of variables with significant differences among stations when 1-
3 way-ANOVAs were performed separately for each port (i.e. 7 in Cagliari, 1 in Heraklion, 0 in El Kantaoui, not
4 shown). The comparison among stations in Cagliari is shown in Table 3 (Scheffé mean multicomparison by
5 stations in Cagliari), where most of the among-station variability was recorded (whole dataset in Table 1S).
6 In Cagliari port, nutrient concentrations were significantly higher and salinity significantly lower in the
7 leisure boat area (C1) than in the other stations. The only exceptions were the intermediate values (not
8 significantly different from those measured in the other stations) found at the port entrance (C5) for
9 salinity and levels of nitrate, phosphate and silicate, as well as at the station hosting the military navy
10 vessels (C2) for the concentration of nitrate. The concentration of total PAHs was significantly higher at the
11 station hosting the military navy vessels (C2) than at the cargo ships station (C4), with intermediate values
12 not significantly different from either in the remaining stations. The overall variance accounted for by PAHs
13 model was 18% of the total variance, as shown by the lower coefficient of determination of PAHs ($r^2=0.18$,
14 Table 3) as compared to the other variables ($r^2>0.70$, Table 3). In the Heraklion port, the only difference
15 found among stations was for silicate, with a coefficient of determination $r^2=0.69$. Silicate concentration in
16 Heraklion was significantly higher in the cargo ship area (H4) than at the port entrance (H7), while values
17 obtained in the other stations were intermediate and not significantly different from either.

20 **3.2. Planktonic biota in the three ports**

21
22 The highest bacterial densities (around 10^7 cell mL⁻¹) were found in Cagliari port in May at all stations (Table
23 2b, Table 1S) and in the three sampling periods at the station hosting leisure boats (C1). On the contrary,
24 the lowest bacterial abundances (around 10^6 cell mL⁻¹) were detected in El Kantaoui in February (all
25 stations) and in Heraklion in September (at the port entrance, H7). The total bacterial counts were
26 significantly higher in Cagliari than in the other two ports (Table 3). No significant differences were found
27 between El Kantaoui and Heraklion ports.

28
29 Considering the overall abundance of phytoplankton, Cagliari port had the significantly highest
30 phytoplankton density at all (total, diatoms and “other phytoplankton”; Table 2b, Table 3), whereas no
31 significant differences resulted for the other two ports. Heraklion was the port with the highest abundances
32 of coccolithophores, which at El Kantaoui presented the lowest values with a significant difference between
33 the two ports; intermediate densities were found in Cagliari (Table 2b, Table 3). The abundances of total
34 phytoplankton showed a general increasing trend in the three ports from the lowest values in February
35 (around 10 cell mL⁻¹) to the maxima recorded in May (El Kantaoui) and September (Cagliari and Heraklion),
36 when phytoplankton reached densities over 10^3 cell mL⁻¹ in Cagliari and El Kantaoui, and 10^2 cell mL⁻¹
37 in Heraklion (Table 2b). Overall, diatoms were the dominant class representing on average more than the 80%
38 of the total phytoplankton in September in the three ports, in May in Cagliari and El Kantaoui, and in
39 February exclusively in Cagliari at stations C1 and C2 (Table 2b, Table 1S). Indeed, in February diatoms and
40 dinoflagellates showed their lowest contribution, when coccolithophores dominated in Heraklion (all
41 stations) and Cagliari (excluding C1 and C2) and “other phytoplankton” dominated in El Kantaoui
42 (particularly cryptophytes) and Cagliari (particularly cryptophytes, freshwater chlorophytes and
43 cyanobacteria). In El Kantaoui (May), Cagliari (May and September) and in Heraklion (September, to lesser
44 extent) diatom blooms with different taxonomic contributions were responsible for the highest densities of
45 total phytoplankton. Dinoflagellates prevailed in Heraklion in May, together with a lower contribution of
46 “other phytoplankton” (cryptophytes and chlorophytes).

47
48 Zooplankton mean abundances ranged between 19 ind m⁻³ in El Kantaoui in February and 941 ind m⁻³ in
49 Heraklion in the same month (Table 2b). Peaks of 2,257 ind m⁻³ in El Kantaoui (September, station E3),
50 2,206 ind m⁻³ in Heraklion (February, Station H3) and 1,542 ind m⁻³ in Cagliari (February, station C3) (Table
51 2S) were recorded. In Heraklion and Cagliari, the highest densities were mostly found in February at the
52 stations hosting passenger and cargo ships (C3 and C4, H3 and H4) and the lowest at the stations hosting
53 leisure boats (C1, H1) in the three sampling periods (Figure 2a, Table 2S). A different pattern was identified
54 in El Kantaoui, where the lowest densities were observed in February and the highest in September (Table
55 2b, Figure 2a).

1 Overall, 39 zooplankton groups were recognised (Table 2b). The most abundant were copepods (67% of the
2 total of all samples), followed by appendicularians (12%) and cladocerans (7%), all belonging to
3 holoplankton (Figure 2a). Meroplankton constituted 12% of the total zooplankton in all the analysed
4 samples, with barnacle nauplii (Cirripedia) contributing with 5%, polychaetes with 3% (mainly spionid
5 larvae) and gastropod larvae and hydromedusae with 1% each.

6 Concerning holoplankton, calanoid copepods represented the 60% of the total zooplankton, with the genus
7 *Acartia* accounting for 44% of the total zooplankton and 67% of the copepods (Figure 2b). Other common
8 calanoid genera identified in the three ports were *Isias* (14% of copepods) very abundant in February in
9 Heraklion, *Paracalanus*, *Parvocalanus*, *Clausocalanus* and *Calocalanus* (reported here as other Calanoida,
10 together with other less abundant or non-identified calanoids, 6% of copepods) and *Centropages* mostly
11 ubiquitous with a 4%. The genus *Acartia* was mostly present in February in Heraklion port, where it formed
12 swarms, declining in May and September. *Acartia* was also observed in Cagliari, mainly in February and May
13 at stations C4 and C5, and in El Kantaoui, mainly in September at the inner stations E1 and E2 (Figure 2b).
14 Cyclopoid copepods were 7% of the copepods, with the genus *Oithona* as the most represented (5%, Figure
15 2b) and very low percentages of Corycaeidae including *Corycaeus*, *Oncoea*, *Farranula* and *Copilia* (3%), the
16 last one exclusively in Heraklion. Harpacticoid copepods were recorded with 2% (Figure 2b), mostly
17 *Diarthrodes* (only at El Kantaoui) and *Euterpina*. The other copepods are included in the group “others”
18 (Figure 2b) and represented less than 1% of the copepods.

19 Appendicularians were the second holoplanktonic dominant taxon, mainly represented by the genus
20 *Oikopleura*, mostly present in Cagliari in February, and in Heraklion and El Kantaoui in September (Figure
21 2a). Cladocerans were observed with all the three genera known for the Mediterranean: *Podon*, *Evadne*
22 and *Penilia*. *Podon* was mostly present in Cagliari in February, with decreasing densities in May and
23 September, whereas *Evadne* was the main genus in Heraklion in September. In El Kantaoui, all the three
24 cladoceran genera occurred with low abundances during the three sampling periods, except for the
25 numerically dominant cladoceran *Penilia* in September.

26 The results of the 2-way-ANOVAs, performed with port and sampling period as main effects on the
27 abundances of total bacteria, total and main groups of phytoplankton, and total zooplankton (Table 3),
28 showed that the variability of the biotic components among ports was higher than the variability among
29 sampling periods, since five of the seven variables presented significant differences among ports (highest
30 abundances in Cagliari for the majority of the variables) and only two among periods (highest abundances
31 of dinoflagellates in May and coccolitophores in February). No significant differences were observed for
32 total zooplankton at all.

33 3.3. Linking abiotic conditions and planktonic elements by factor analysis

34 The combined effects of the variables were analysed through factor analysis and the first four factors
35 explained 57% of the whole data variability (Table 4). Each factor includes a different combination of
36 variables with coefficients, which reflects different sources of variability in the studied data. The
37 coefficients of each variable are relevant for the interpretation of the factor itself (Table 4). The values of
38 the four factors calculated for each station and sampling period in the three investigated ports are reported
39 in Figure 3.

40 The first factor (Factor1, Table 4) accounted for 24% of the overall data variability. It is a bipolar factor
41 showing in its positive pole a strong positive correlation (high positive coefficients) among DO,
42 concentrations of nutrients (NH₄, NO₂, NO₃, PO₄, SiO₂), POC, chlorophyll-*a*, total bacterial counts, as well as
43 abundances of diatoms, “other phytoplankton” and zooplanktonic cladocerans and barnacles nauplii
44 (Cirripedia), while a weaker correlation (mid positive coefficients) was found with the abundances of
45 appendicularians and decapod larvae. These variables were negatively correlated with those in the negative
46 pole, strongly (high negative coefficient) with salinity and weakly (mid negative coefficient) with the
47 abundance of ichthyoplankton (Table 4). Factor1 differentiated Cagliari from the other two ports, with
48 higher values for Cagliari than El Kantaoui and Heraklion in all stations and sampling periods, as result of
49 lower salinity, higher nutrient levels and higher plankton abundances (Figure 3a). In time and at station
50 scale, the Cagliari port presented higher values of Factor1 in May at all stations and in the three sampling
51 periods at the stations hosting leisure boats (C1) and military navy vessels (C2) as well as at the port
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entrance (C5). In the ports of Heraklion and El Kantaoui, Factor1 values were rather similar among different stations in the same sampling period, showing an increasing trend with time.

The second factor (Factor2, Table 4) accounted for a further 15% of the remaining data variability. It reflected the abundances of most zooplankton taxa (both holoplanktonic and meroplanktonic), more specifically, higher coefficients were found for copepods (harpacticoids, cyclopoids, calanoids and monstrilloids), hydromedusae, chaetognats, appendicularians, molluscs (both gastropods and bivalves) and polychaetes, while lower coefficient values were found for barnacle nauplii. Among the investigated ports, the variability range of Factor2 was wider and the differences through time were larger for El Kantaoui (Figure 3b), with lower values in February and higher in September, mainly at the port entrance (E3). On the contrary, the variability ranges were rather similar and narrow in Cagliari and Heraklion, with higher differences between ports than sampling periods. In the two larger ports, the Factor2 generally increased from the inner leisure boat area towards the port entrance (in Heraklion up to H4, not reaching H5 and H7) and at each station from February to May and September (Figure 3b). Overall, Factor2 differentiated zooplanktonic communities between El Kantaoui and the other two ports, but also among different stations within each port and through time, representing the second main source of variability in the analysed data. The third factor (Factor3, Table 4) accounted for a further 11% of the remaining data variability. It can be described as a bipolar factor showing negative correlations between concentrations of nitrate and silicate, abundances of coccolithophores and cyclopoid copepods on the positive pole, and temperature, abundances of dinoflagellates, diatoms and platyhelminth larvae on the negative pole. A similar trend was recognized in Cagliari and Heraklion (Figure 3c), with higher values of Factor3 in February (as result of the maxima of coccolithophores and the minima of diatoms and dinoflagellates) and lower values in May and September (as result of the maxima of dinoflagellates in May and the maxima of diatoms in May and September). On the contrary, a different trend was found in El Kantaoui, with the lowest values of Factor3 in May, corresponding to a synchronous variation of the planktonic taxa included in the factor at the negative pole at the three sampling stations. In all ports and stations, Factor3 differentiated samples collected in winter (February) from samples collected during the warmer months (May and September, Figure 3c). Moreover, in February the Factor3 values discriminate between the two large ports (Cagliari and Heraklion with higher values of Factor3) and the marina of El Kantaoui (lower values of Factor3).

The fourth factor (Factor4, Table 4) accounted for a further 7% of the remaining data variability. It is a bipolar factor showing negative correlation between the abundances of siphonophores, echinoderms larvae and ichthyoplankton in the positive pole, and concentrations of AHs and PAHs as well as abundances of amphipods in the negative pole. Moreover Factor4 resulted significantly correlated with water depth (Pearsons $r^2 = 0.34$, $p < 0.05$; $r^2 = 0.44$, $p < 0.05$ excluding the winter Cagliari samples). The shallow El Kantaoui and the deep Heraklion ports presented opposite patterns, while the intermediate-depth Cagliari port was similar to El Kantaoui in February and to Heraklion in May and September (Figure 3d). In El Kantaoui, the coefficients of Factor4 were lower as compared to the other ports at all stations and in all sampling periods (Figure 3d), with the exception of the samples collected in winter in Cagliari at the stations hosting the military navy vessels (C2), passenger (C3) and cargo ships (C4) (Figure 3d). The generally lower Factor4 values in El Kantaoui were the result of higher levels of AHs and amphipods and lower (or zero) levels of the three positive coefficient variables (i.e. abundances of siphonophores, echinoderms larvae and ichthyoplankton). An opposite situation was found in Heraklion, where the coefficients of Factor4 were higher than in the other two ports in almost all the sampling stations and periods, as related to higher abundances of ichthyoplankton, echinoderm larvae and siphonophores and lower levels of AHs, PAHs and amphipods (Figure 3d). In Cagliari port, the coefficients of Factor4 presented lower values in February (more similar to those measured in El Kantaoui) than in the other sampling periods, when higher levels of hydrocarbons and amphipods were measured in the port waters. On the contrary, higher Factor4 values (more similar to those observed in Heraklion) were found at all the stations of the Cagliari port in May, when concentrations of PAHs were lower and echinoderm larvae were more abundant (Table 2b).

4. Discussion

4.1. Differences in freshwater inputs and nutrient loading

1 At the space and time scales of the present study, the main variability occurred among ports as it was
2 identified by Factor1. Overall, statistical analyses cogently differentiated the port of Cagliari from the port
3 of Heraklion and the marina of El Kantaoui (Table 3 and Figure 3a). The specificity of the Cagliari port can be
4 attributed to the input of inland brackish waters into the port area, a hydrological feature that does not
5 occur in the other two harbours. The Natural Protected Area of Molentargius is a system of coastal lagoons,
6 ponds and saltworks protected by the RAMSAR Convention. Within this system, the San Bartolomeo canal
7 (Figure 1b) receives water from the brackish lagoon “Stagno del Molentargius” and collects treated
8 wastewaters from the surrounding Municipalities, discharging brackish waters mainly into the leisure boat
9 area (C1) of the Cagliari port (Figure 1a, b). A plume of this water likely extends to cover the area of the
10 port entrance (C5) providing a distinct water profile within the port. The effects were mainly visible in
11 February in the area hosting leisure boats (C1, minimum salinity 28‰) and at the port entrance (C5,
12 minimum salinity 30‰). This discharge affected not only salinity but also nitrate, phosphate and silicate (at
13 the negative pole of Factor1, Table 4) with high levels of the three nutrients at stations C1 and C5, and high
14 level of nitrate at station C2 (1-way-ANOVA, Table 3). In the proximity of the quays for the anchoring of
15 cargo ships (C4), the Santa Gilla canal drains wastewaters of urban runoff into the port area (Figure 1b).
16 Despite this freshwater inlet, the station C4 seemed to be marginally influenced, probably due to a low flow
17 rate and/or the quality of the discharged waters. Finally, the sector of the Cagliari port hosting passenger
18 ships (C3), more distantly located from the two discharges of inland waters, was the least affected area of
19 the port (highest salinity and lowest nutrient levels during all the studied periods).

20 The brackish water input of the Molentargius lagoon not only drains into the Cagliari port high levels of
21 inorganic nutrients and organic matter (Massi et al., 2019), but also plankton. Freshwater chlorophytes,
22 cyanobacteria (namely “other phytoplankton”, Table 3) and the freshwater pennate diatom *Tabellaria*
23 *fenestrata*, typical of eutrophic waters, were exclusively found in Cagliari among the studied ports,
24 particularly at stations C1 and the nearby C2. The high trophic level of waters in the Cagliari port was also
25 likely responsible for the development of diatom blooms and the proliferation of bacteria, which affected
26 station C1 (and also C2 for diatoms) in the three studied periods, and influenced all the stations in May,
27 probably due to persistent effects of nutrient rich brackish waters from the Molentargius lagoon after the
28 winter-spring rainy period. More specifically, these high trophic conditions promoted the May-September
29 blooms of typical coastal fast-growing and individually small diatoms, as *Skeletonema pseudocostatum* and
30 *Thalassiosira pseudonana*. Indeed, *Skeletonema* species are well known blooming diatoms in coastal and
31 estuarine Mediterranean waters (Moncheva et al., 2001; Kooistra et al., 2008; Abboud-Abi Saab et al.,
32 2008), while *T. pseudonana*, typical of coastal and brackish environments, is favoured by high temperatures
33 (Hegseth and Sakshaug, 1983).

34 Concerning zooplankton, three variables of the positive pole of Factor1 highlighted the association of the
35 eurihaline and neritic cladoceran *Podon* (Factor1, Table 4) with crustacean larvae and the appendicularian
36 *Oikopleura dioica*. The first two groups are known in disturbed and shallow areas subjected to river
37 discharges (Christou et al., 1995) and the latter depends on chlorophyll and temperature for its
38 development (Harris et al., 2005). Specifically, the genus *Podon* is described as a raptorial feeder on big
39 particles, such as microzooplankton and large phytoplankton (Jagger et al., 1988) that are abundant in the
40 water column in eutrophicated areas with high nutrient loading. Accordingly, this taxon was present in
41 Cagliari with the highest densities during all the sampling periods, constituting one of the distinct features
42 of the Cagliari port among the studied harbours.

43 As compared to the Cagliari port, more saline waters with lower levels of nutrients, bacteria, phytoplankton
44 and lower abundance of zooplankton with positive coefficients in Factor1 prevailed in Heraklion and El
45 Kantaoui ports in all the studied periods. A trend is observed in these ports with a mild increase of Factor1
46 from winter to summer (Factor1, Figure 3a), which may be attributed to an increased nutrient loading
47 during the tourist warm season together with increases of some plankton organisms. In El Kantaoui, the
48 diatom blooming in May was found exclusively for *S. pseudocostatum*. As a matter of fact, *Skeletonema*
49 blooms have been observed in the port waters of Malta (De Bono, 2001/2002; Nuccio, unpublished data)
50 and other southern Mediterranean ports (Abdel-Halim and Khairy, 2007; Heneash et al., 2015). In
51 September, a contribution to the aforementioned increasing trend was given by the large occurrence in El
52 Kantaoui port of the filter-feeder cladoceran *Penilia avirostris*, described as more typical of warm season
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1 and waters (Margaritora, 2010), and by the high abundances in Heraklion of Cirripedia and *Oikopleura*
2 (Figure 2a), the latter probably dependent on food availability as one of the most critical limiting factors
3 (Tomita et al., 2003).

4 The further variable contributing to the different patterns observed between Cagliari port, on one side, and
5 El Kantaoui and Heraklion, on the other side, is the higher density of ichthyoplankton that occurred in the
6 last two ports. Ichthyoplankton seemed more connected with open sea processes and was indeed mainly
7 observed at the outer stations of El Kantaoui and Heraklion ports. Its low abundance in Cagliari might be
8 related with the freshwater inputs and/or advection from neighbouring areas.

9 From the point of view of the relationships among the biotic components highlighted by Factor1, bacteria
10 and phytoplankton directly benefit of nutrient loading whereas the zooplankton organisms, like
11 cladocerans, barnacle larvae and appendicularians, take advantage of both suspended particles and high
12 phytoplankton abundances even if bacterioplankton should not be excluded (Bernàt et al., 1994). Among
13 abiotic parameters, the three biotic components are also strongly influenced by salinity. At the highest
14 trophic level in the studied sites (Rice and Williamson, 1970), decapod larvae benefit of the presence of the
15 other planktonic groups.
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18 **4.2. Zooplankton variability**

19 Besides the seasonality of the different zooplanktonic species, local processes as tidal oscillations, bottom
20 processes, wind stress and turbulence causing turbidity and/or low hydrodynamics are considered the main
21 drivers enhancing or reducing abundances, and structuring zooplanktonic communities and their biological
22 interactions in harbours and bays (Dawson and Pieper, 1993). Parallel considerations may be made for the
23 present study in which Factor2 shows a non-homogeneous distribution of the relevant zooplanktonic taxa
24 and their variability among stations within each port and with season (Factor2, Table 4, Figure 3b).

25 In both large ports of this study (Cagliari and Heraklion), the wide connection between the port area and
26 the open sea through their wide port entrance, the relatively deep bathymetry and the wide area between
27 docks and breakwaters (C3-C5 and H3-H5, Figure 1) allow a rather free water penetration from the open
28 sea. These conditions result in a higher abundance of specific groups of zooplankton (as shown by Factor2,
29 Table 4) at the port entrance (C5 and H7) as compared to the respective shallower and more protected
30 areas (C1, C2 and H1, Figure 3b), where open sea water penetration is more difficult. Furthermore, in C1
31 and C2 a disturbance element for zooplankton grazing could have been the presence of cyanobacteria
32 (included in "other phytoplankton" and found exclusively in Cagliari), for their eventual toxin production
33 and poor manageability by zooplankton (Hogfors et al., 2014, and references therein). In contrast, the
34 shallow and narrow port entrance in the smaller El Kantaoui port is likely to reduce water penetration
35 inside the marina (Figure 1, Table 1). Despite plankton advection can be supposed to influence plankton
36 changes in stations close to the port entrance, investigation of the phenomenon was out of the aim of the
37 study and can not be achieved through the present sampling protocol.
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40 In the El Kantaoui marina in February, the relatively higher winter water mixing and the zooplankton
41 seasonality would explain the similar low zooplankton abundance in the port entrance (E3) and
42 intermediate area (E2), and the even lower abundance in the leisure boat station (E1) (Figure 3b); this
43 spatial difference suggests that the open sea water should have penetrated into the port up to station E2.
44 From May to September, the low hydrodynamism seems to have favoured a local zooplankton
45 development inside the port (positive value of Factor2 at station E1, Figure 3b). Indeed, very still water was
46 documented during the whole sampling campaign in September, which could account for the much higher
47 zooplankton abundance found at the port entrance (E3) than at the inner stations (peak of Factor2, Figure
48 3b). It is worth noting that at station E3 the open sea taxa (e.g. hydromedusae, appendicularians and
49 gastropod larvae) were observed in higher abundances compared to the densities in the other stations,
50 while opportunistic taxa (e.g. calanoids and barnacle larvae) were more abundant in the inner station E1
51 (Table 2S).
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54 Among copepod genera, harpacticoids, cyclopoids and calanoids had a considerable weight in the
55 zooplankton communities as indicated by Factor2 (Table 4). Among cyclopoids, the most abundant taxon
56 was *Oithona*, which represents an opportunistic zooplankton, belongs to the coastal neritic or shallower
57 waters (Williams and Muxagata, 2006) and is known to feed on a wide range of particles, such as organic
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1 matter and phytoplankton of small size (Lampitt and Gamble, 1982; Turner, 2004). The calanoid *Acartia*
2 was the most conspicuous copepod at the three ports (Figure 2b). It is a well-known swarming genus (Ueda
3 et al, 1983; Santu et al 2016) that may take advantage of coastal areas with suitable conditions like
4 sheltered bays rich in nutrients, but also port areas (Siokou-Frangou et al., 1995; Belmonte et al., 2018,
5 Vidjak et al., 2018). In literature, it has been reported that much of *Acartia* seasonality depends on
6 temperature, salinity and hydrology and eventually oxygen or chlorophyll-*a* (proxy of phytoplankton)
7 (Siokou-Frangou et al., 1998; Kang, 2011). The *Acartia* abundances observed in this study (Figure 2b)
8 apparently followed the trends typical of naturally enclosed coastal areas with the booster effect of
9 nutrient abundance that favoured phytoplankton blooms (Siokou-Frangou and Papathanassiou, 1991). An
10 inverse relationship between the relative quantities of phytoplankton and zooplankton have been observed
11 by many authors and were explained with zooplankton grazing, animal exclusion in phytoplankton patches,
12 or the different reproduction rates of vegetal and animal populations (Cattani and Corni, 1992). In this
13 study, the sampling periodicity was not frequent enough to define a seasonal relationship between phyto-
14 and zooplankton. Nevertheless, in El Kantaoui high abundances of calanoids (*Acartia*) were observed in
15 May and September along with diminishing abundances of diatoms, most likely ascribable to the action of
16 copepods grazing (Ryther and Sanders, 1980) favoured by the sheltered waters of the inner stations E1 and
17 E2 (Table 2S).
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20 Predators like hydromedusae and chaetognaths that have restricted ranges of tolerance to pollution and
21 variation of environmental parameters, were present in the outer port stations, where advection can affect
22 the plankton community structure (Bracco et al., 2009). On the other hand other more opportunistic or
23 tolerant taxa like polychaetes, bivalves and appendicularians seemed to benefit of the port environment
24 mainly occupying the intermediate area of the port, but not the inner one (Table 2S). A similar pattern was
25 described by Siokou-Frangou and Papathanassiou (1991) that resumed how the most opportunistic species
26 populate the most disturbed areas characterized by the absence of carnivorous species.
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28 With the exception of the cladoceran *Podon* sp. strongly contributing to Factor1, most zooplankton taxa
29 have very low coefficients in Factor1 and Factor3 (Table 4). Therefore, the zooplanktonic communities are
30 marginally influenced by those variables playing a crucial role for phyto- (Factor1 and 3) and bacterio-
31 plankton (Factor1). Specifically, copepods and hydromedusae (that have high coefficients in Factor2, Table
32 4) depend on interspecific interactions as observed by Pesticó et al. (2018) and polychaete and bivalve
33 larvae are interconnected to the other taxa, even if belonging to the lower levels of the food web.
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36 **4.3. Seasonality and water turbulence**

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38 Seasonal differences among ports were highlighted by Factor3 (Table 4, Figure 3c) with rather opposite
39 trends between Cagliari and Heraklion on one side, and El Kantaoui, on the other, reasonably related to
40 changes in water temperature and seasonal dominance of specific phytoplankton groups (diatoms,
41 dinoflagellates, coccolithophores). Winter conditions of major wind stress, water mixing and turbulence,
42 likely contributed to particle re-suspension from the bottom and could favour higher concentrations of
43 nitrate and silicate in the water column (Table 2a). This phenomenon was more evident in the two larger
44 ports of Cagliari and Heraklion as compared to the marina of El Kantaoui (Figure 3c). Under these
45 environmental conditions, seasonal changes in phytoplankton composition and cyclopoids density have an
46 important role in accounting the variance in Factor3 (Table 3, Table 4).
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48 Coccolithophores resulted generally more abundant in winter, most noticeably in the two larger ports and at
49 the stations most exposed to seawater fluxes, likely shaped by currents and greatly reduced during the
50 warmer periods (Table 2b). This taxon was demonstrated to constitute a large part of the nano-
51 phytoplankton fraction in the Mediterranean Sea, recorded mainly in autumn and winter in south-eastern
52 waters (Siokou-Frangou et al., 2010). Moreover, their presence in ports was linked to seawater flux by
53 Massi et al. (2019). Compared to coccolithophores, in Cagliari and Heraklion diatoms and dinoflagellates
54 had an opposite trend, increasing their abundances from February to May and September (Table 2b).
55 Moreover, the observed negative correlation of diatoms with silicate concentration at opposite poles of
56 Factor3 (Table 4) may depend on the feeding of diatoms consuming the nutrient. In El Kantaoui a different
57 pattern of Factor3 was evident for the absence of coccolithophores in winter, except for a very scarce
58 density at the port entrance, as well as the high diatom bloom and the increment of dinoflagellates in May.
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1 These blooming diatoms are typical of shallow waters over a wide range of temperatures for their
2 opportunistic features of exploiting nutrients and organic matter (Carstensen et al., 2015).

3 Concerning the zooplanktonic components at the Factor3 positive pole, almost all the cyclopoids belonged
4 to the genus *Oithona*, which are typical of neritic areas and enclosed systems, frequently associated with
5 nutrients and presenting very flexible relationships with temperature (Mazzocchi and Ribeira D'Alcalà,
6 1995; Calbet et al., 2001; Dvoretzky and Dvoretzky, 2015; Ben Ltaief et al., 2015; Wang et al., 2017). *O.*
7 *plumifera* (Wang et al., 2017) and *O. similis* (Dvoretzky and Dvoretzky, 2015) have been demonstrated to be
8 negatively influenced by water temperature, while *O. nana* seasonality was described as changing also
9 based on latitude (Williams and Muxagata, 2006). Therefore it is not to be excluded the same response in
10 Cagliari and Heraklion ports where cyclopoids contributed to Factor3 in February with respect to the
11 opposite trend observed in El Kantaoui, characterised by increasing concentrations of *Oithona* and other
12 cyclopoids from February and May to September. Moreover, as for the other planktonic components, in
13 rich and favourable conditions the environmental variables affecting life cycles of specific zooplanktonic
14 taxa may shift to a different hierarchy driven by local dynamics.

15 Factor3 highlighted the importance of the variation of abundances of platyhelminthes, a taxon that showed
16 similar pattern of diatoms and dinoflagellates and was mostly found in calm and warmer conditions.
17 Indeed, the taxon was mainly represented by Müller larvae and was collected mostly in May at El Kantaoui.
18 This result may depend on the species local life cycle and feeding preferences (Rawlinson, 2014). As a
19 matter of fact, a high concentration of Müller larvae was as well found during a further sampling in June
20 2015 in the same marina (Rossano, unpublished data).

21 22 23 24 **4.4 Impacts of anthropogenic activities**

25 The last factor explaining the variability in the analysed biotic and abiotic parameters is related to
26 anthropogenic activities as the main sources of PAHs and AHs contaminating port waters (Factor4, Table 4).
27 Pyrogenic emission sources associated to the incomplete combustion of fuels and biomasses (e.g. fuel
28 combustion in engines) are the main origin of PAHs in the three studied ports (Vitali et al., 2019) in line with
29 several other Mediterranean harbours (Merhaby et al., 2015; Schintu et al., 2015). In addition, accidental
30 oil spills and leakages of refined oil products (e.g. diesel, lubricating oils) are anthropogenic sources of
31 petrogenic PAHs and AHs entering the port waters, even if AHs may also derive from natural origins, such
32 as biomass of marine microorganisms (i.e. phytoplankton, algae and bacteria) and transfer of terrestrial
33 plant detritus from the land into the sea (Head et al., 2006; Mandalakis et al., 2014; Chatzinikolaou et al.,
34 2018; Vitali et al., 2019).

35 The concentrations of PAHs in surface waters of the three ports (24 - 336 ng L⁻¹, Table 1S) were within the
36 ranges previously recorded in Mediterranean open sea (10 - 30 ng L⁻¹ in North Aegean Sea, Abdulla and
37 Linden, 2008) and coastal waters (20 - 40,000 ng L⁻¹ in Turkish coasts, Abdulla and Linden, 2008) and the
38 levels were well below the concentrations leading to 50% mortality (300,000 - 2,500,000 ng L⁻¹, Kennish,
39 1998) or producing chronic effects on most marine organisms (50,000 - 150,000 ng L⁻¹, ANZECC, 1999).
40 Therefore, the concentration of PAHs does not seem high enough to have a strong impact on the
41 zooplankton, as it was also established in the same sites by Chatzinikolaou et al. (2018) for benthic
42 macrofauna and by Tamburini et al. (2020) for benthic prokaryotes.

43 Among the investigated harbours, the Cagliari port exhibited the highest concentrations of PAHs in surface
44 water, according to the levels of PAH contamination in sediments (Vitali et al., 2019). Nevertheless,
45 sediment levels of PAHs in the three studied ports spanned within a wider range (25 - 49,000 ng g⁻¹) as
46 compared to concentrations in the water column (Vitali et al., 2019). Hydrocarbons, and particularly PAHs,
47 tend to associate with particulate matter due to their low water solubility, sinking to the bottom and
48 accumulating in sediments over time (Readman et al., 2002; Zakaria et al., 2002). This explains why their
49 concentrations in the studied sites were considerably higher in sediments than in the overlying water
50 column, as frequently observed in literature (Abdulla and Linden, 2008). On the other hand, hydrocarbon
51 degrading bacteria were found to be abundant in surface waters at the three studied ports, where they
52 seem to be involved in the fate of hydrocarbons in the water column (Bullita et al., 2014; Bullita et al.,
53 2016). Indeed, decreasing concentrations from winter to the warm periods was evident in Cagliari both for
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AHs and PAHs and in El Kantaoui for PAHs, which may be at least partially ascribed to the low degradation rate at winter temperature (Head, 2006; Bullita et al., 2016).

In shallow waters, besides the direct inputs of hydrocarbons by anthropogenic activities, sediment re-suspension occurs through different processes (e.g. navigating vessels, turbulence due to storms, wind and winter water mixing, tides) and may cause hydrocarbon re-mobilization into the water column (Roberts, 2012). In the investigated ports, this phenomenon seems to occur in the small marina of El Kantaoui, characterized by a low bathymetry in the whole port area, as well as at the shallow stations of the two bigger ports of Cagliari (C2) and Heraklion (H1) (Table 1), where low Factor4 values and high hydrocarbon levels were mostly found (Figure 3d). More specifically, benthos samples collected at station C2 in Cagliari revealed a black fatty substance that glued the sediments making the sieving process very difficult (Chatzinikolau et al., 2018; Rossano C., personal field observations). Indeed, the levels of PAHs in sediments of station C2 (49,000 ng g⁻¹) resulted one order of magnitude higher than the concentrations found in the other sectors of the Cagliari port and the highest among the three investigated harbours (Vitali et al., 2019); consistently, the highest concentration of PAHs in surface water were found in the present study at station C2 (336 ng L⁻¹ in February). In Cagliari, burning of coal and biomass are the main source of PAHs entering the port water by atmospheric deposition and street run-offs emitted from the adjacent city (Vitali et al., 2019), a pollution usually characterised by recalcitrant compounds with a long-term persistence in marine environments (Yunker et al., 2002; Duran and Cravo-Laureau, 2016). Therefore, the local high contaminations of PAHs at the station C2 could be reasonably attributed to the presence in the past of a water drainage channel, which collected city run-offs to this shallow area of the Cagliari port channel (RAS-ARDIS local Authorities, personal communication, 2015).

In line with the close interconnection between water column and benthos, Factor4 was significantly correlated with water depth, a parameter that is directly linked to the specific anthropogenic activities operated in each port sector (Table 1). Therefore, water depth seems to be an important descriptor of the on-going processes in ports. An exception to this trend was the negative Factor4 values found in February in the Cagliari port not only at the shallow water station C2, but also in the deep water stations (Figure 3d), which may be reasonably explained by an increased water turbulence and consequent sediment re-suspension under the particularly windy conditions during the winter sampling.

Concerning fauna at the shallow water stations, benthic amphipods (the only biotic component at the negative pole of the Factor4) were found in the water column likely because of their increased mobility, or due to the particle re-suspension and consequent hydrocarbon mobilisation. On the opposite, the positive pole organisms (siphonophores, echinoderms, ichthyoplankton) prevailed in Cagliari and Heraklion ports and were more abundant in the deep stations (Figure 3d, high values in Heraklion in H3-H5 and in Cagliari in C3 and C4), likely related to local variation in community composition and more favourable conditions in waters with less sediment re-suspension.

Conclusions

This study assessed for the first time the three components of the planktonic biota (bacterio-, phyto- and zooplankton) in different ports at a large spatial scale (i.e. Mediterranean basin) and compare them with respect to environmental variables and anthropogenic inputs. The description of the planktonic biota in the three ports (aims i and ii) revealed generally abundant and complex communities with different relationships with water abiotic components (aim ii) in the three ports. A high influence of brackish eutrophic water discharge was highlighted in the port of Cagliari causing high abundances of bacterioplankton and spring-summer blooms of phytoplankton. Unlike the other two components of plankton, many zooplankton taxa resulted influenced by complex interspecific interactions, but in general, nutrient loading played an important role in their distribution both among different ports and within ports. Seasonality was also clear depending more on meteorological factors (temperatures, winds and storms) than on human activities (tourist season). The studied ports seem to guarantee rich ecosystems, favouring the growth of zooplanktonic communities with a specific periodicity (aim iii) obviously facilitating those opportunistic species that are already known in nutrient-rich coastal areas. The different port activities apparently did not affect planktonic networks, which were more linked to inland loading and seasonal conditions. Hydrocarbon pollution did not specifically affect phyto- and zooplankton components and

1 seems to be controlled by degradation activities of bacterioplankton. The described ecosystem variability
2 emphasizes the importance of the relative contribution of inland, port and marine waters and suggest
3 eventual critical points to take into account in view of a sustainable management of port areas. It is
4 therefore suggested to include the planktonic community as a whole (bacterioplankton, phytoplankton and
5 zooplankton), in the monitoring programs of port areas to control the quality of these heavily modified
6 water bodies and their impacts on coastal waters.

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Table 1. Location of the sampling stations in the three Mediterranean ports (Cagliari, Sardinia, Italy; El Kantaoui, Sousse, Tunisia; Heraklion, Crete, Greece) and main uses of the sampling stations within each port (Chatzinikolaou et al., 2018; Massi et al., 2019; Vitali et al, 2019).

| Station | Latitude | Longitude | Water Depth (m) | Station use |
|--------------------|---------------|---------------|-----------------|--|
| Cagliari | | | | |
| C1 | 39°12'11.40"N | 9° 7'24.12"E | 7.8 | Leisure - small boats |
| C2 | 39°12'20.46"N | 9° 7'15.06"E | 4.5 | Intermediate (military navy vessels) |
| C3 | 39°12'27.12"N | 9° 6'46.44"E | 8.3 | Passenger ships |
| C4 | 39°12'25.98"N | 9° 6'18.60"E | 13.5 | Cargo ships |
| C5 | 39°11'52.94"N | 9° 6'41.10"E | 11.4 | Port entrance (oriented to south) |
| El Kantaoui | | | | |
| E1 | 35°53'38.64"N | 10°35'53.16"E | 2.5 | Leisure - small boats |
| E2 | 35°53'34.44"N | 10°35'58.92"E | 4.0 | Intermediate (fuel station) |
| E3 | 35°53'34.65"N | 10°36'4.44"E | 3.2 | Port entrance (oriented to south-east) |
| Heraklion | | | | |
| H1 | 35°20'36.32"N | 25° 8'9.93"E | 3.7 | Leisure - small boats |
| H3 | 35°20'44.70"N | 25° 8'40.87"E | 19.5 | Passenger ships |
| H4 | 35°20'42.70"N | 25° 8'52.28"E | 10.5 | Cargo ships |
| H5 | 35°20'48.72"N | 25° 9'7.94"E | 19.0 | Shipyard |
| H7 | 35°20'50.82"N | 25° 9'17.88"E | 7.0 | Port entrance (oriented to east) |

Table 2a. Mean values of physical and chemical variables by port and sampling period in the three Mediterranean ports

| Variable | Unit | Cagliari | | | El Kantaoui | | | Heraklion | | |
|-----------------------|--------------------|----------|-------|-------|-------------|-------|-------|-----------|-------|-------|
| | | Feb | May | Sep | Feb | May | Sep | Feb | May | Sep |
| ENVIRONMENT | | | | | | | | | | |
| Temperature | °C | 10.3 | 21.0 | 22.3 | 11.7 | 23.7 | 26.5 | 14.6 | 21.0 | 25.3 |
| Salinity | ‰ | 33.2 | 33.2 | 34.2 | 37.0 | 36.9 | 36.1 | 38.2 | 38.3 | 37.3 |
| DO | mg L ⁻¹ | 9.3 | 11.5 | 7.5 | 8.7 | 6.3 | 4.5 | 7.8 | 7.0 | 6.7 |
| pH | | 8.1 | 8.3 | 8.4 | 8.2 | 8.2 | 8.2 | 8.2 | 8.2 | 8.4 |
| Ammonia | μM | 10.9 | 14.9 | 12.6 | 0.3 | 0.2 | 2.3 | 0.5 | 0.5 | 0.5 |
| Nitrite | μM | 1.65 | 3.00 | 1.62 | 0.00 | 0.03 | 0.17 | 0.05 | 0.01 | 0.04 |
| Nitrate | μM | 26.7 | 18.5 | 12.9 | 0.1 | 0.7 | 0.6 | 7.7 | 9.8 | 6.9 |
| Phosphate | μM | 1.40 | 3.78 | 3.08 | 0.07 | 0.05 | 0.22 | 0.14 | 0.04 | 0.02 |
| Silicate | μM | 11.52 | 9.93 | 6.29 | 1.39 | 0.39 | 1.81 | 6.26 | 3.74 | 4.44 |
| POC | μg L ⁻¹ | 1,270 | 2,781 | 1,768 | 1,013 | 906 | 1,202 | 1,202 | 1,068 | 1,094 |
| Chlorophyll- <i>a</i> | μg L ⁻¹ | 2.80 | 10.50 | 5.18 | 0.35 | 0.95 | 0.34 | 0.39 | 0.50 | 0.99 |
| AHs | ng L ⁻¹ | 5,257 | 3,888 | 2,720 | 3,133 | 3,867 | 3,196 | 2,340 | 1,969 | 3,031 |
| PAHs | ng L ⁻¹ | 137 | 68 | 43 | 103 | 42 | 37 | 46 | 60 | 67 |

DO: dissolved oxygen; POC: particulate organic carbon; AHs: aliphatic hydrocarbons; PAHs: polycyclic aromatic hydrocarbons.

Table 2b. Mean values of biological variables by port and sampling period in the three Mediterranean ports

| Variable | Unit | Cagliari | | | El Kantaoui | | | Heraklion | | |
|---|------------------------|----------|----------|----------|-------------|----------|--------|-----------|-------|--------|
| | | Feb | May | Sep | Feb | May | Sep | Feb | May | Sep |
| BACTERIA & PHYTOPLANKTON | | | | | | | | | | |
| Total bacterial counts x10 ⁶ | cells mL ⁻¹ | 7.6 | 8.9 | 7.0 | 1.9 | 4.2 | 6.3 | 4.7 | 3.8 | 2.7 |
| Total phytoplankton | cells mL ⁻¹ | 221.00 | 3,813.00 | 4,364.00 | 29.00 | 1,421.00 | 141.00 | 24.00 | 65.00 | 100.00 |
| Diatoms | cells mL ⁻¹ | 171.00 | 3,280.00 | 3,754.00 | 10.10 | 1,338.00 | 115.00 | 1.40 | 11.90 | 73.30 |
| Dinoflagellates | cells mL ⁻¹ | 3.40 | 16.00 | 14.00 | 1.20 | 38.00 | 2.90 | 4.10 | 27.00 | 13.00 |
| Coccolitophores | cells mL ⁻¹ | 6.70 | 0.40 | 1.80 | 0.06 | 0.20 | 0.00 | 14.00 | 3.40 | 1.40 |
| Other phytoplankton | cells mL ⁻¹ | 40.00 | 516.00 | 594.00 | 17.00 | 45.00 | 23.00 | 3.80 | 23.00 | 12.00 |
| ZOOPLANKTON | | | | | | | | | | |
| Total zooplankton | ind m ⁻³ | 540.0 | 475.0 | 128.0 | 19.0 | 140.0 | 910.0 | 941.0 | 112.0 | 322.0 |
| Holoplankton | | | | | | | | | | |
| Proportion of holoplankton | % | 94.4 | 72.7 | 57.8 | 48.9 | 25.9 | 77.6 | 95.6 | 67.2 | 83.0 |
| Hydromedusae | ind m ⁻³ | 5.4 | 3.1 | 4.8 | 0.3 | 0.3 | 45.0 | 0.8 | 3.1 | 3.1 |
| Scyphomedusae | ind m ⁻³ | 7.5 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| Siphonophora | ind m ⁻³ | 0.1 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 2.7 | 0.2 | 0.0 |
| Pteropoda | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.1 | 0.0 | 0.0 | 0.0 | 0.2 | 0.4 |
| Ostracoda | ind m ⁻³ | 0.3 | 0.7 | 0.0 | 0.1 | 0.0 | 1.0 | 0.1 | 0.7 | 0.1 |
| <i>Podon</i> spp. | ind m ⁻³ | 125.5 | 85.4 | 18.3 | 0.3 | 1.1 | 0.0 | 0.1 | 0.0 | 0.0 |
| <i>Evadne</i> sp. | ind m ⁻³ | 0.0 | 0.0 | 0.1 | 0.0 | 0.8 | 11.6 | 0.0 | 0.0 | 9.6 |
| <i>Penilia</i> sp. | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 24.4 | 0.0 | 0.0 | 0.0 |
| <i>Acartia</i> spp. | ind m ⁻³ | 64.4 | 119.6 | 2.4 | 1.1 | 6.3 | 163.0 | 644.1 | 56.3 | 67.0 |
| <i>Centropages</i> spp | ind m ⁻³ | 2.4 | 34.3 | 11.3 | 0.0 | 2.6 | 27.6 | 3.4 | 1.6 | 18.3 |
| <i>Isias</i> sp | ind m ⁻³ | 0.4 | 1.8 | 0.3 | 0.0 | 0.0 | 0.0 | 187.1 | 0.4 | 3.5 |
| other Calanoida | ind m ⁻³ | 24.2 | 28.4 | 3.6 | 2.2 | 2.3 | 98.7 | 27.6 | 6.7 | 17.2 |
| <i>Corycaeus</i> sp. | ind m ⁻³ | 1.4 | 0.0 | 0.4 | 0.0 | 0.0 | 54.3 | 9.6 | 0.6 | 1.1 |
| <i>Oithona</i> spp. | ind m ⁻³ | 41.9 | 13.3 | 2.8 | 2.2 | 1.4 | 82.4 | 13.3 | 2.9 | 17.9 |
| other Cyclopoida | ind m ⁻³ | 0.2 | 0.1 | 0.0 | 0.8 | 0.0 | 1.8 | 1.2 | 0.1 | 0.0 |
| <i>Diarthrodes</i> sp. | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.1 | 1.5 | 64.8 | 0.0 | 0.0 | 0.0 |
| <i>Euterpina</i> sp. | ind m ⁻³ | 4.2 | 3.8 | 2.0 | 0.5 | 1.0 | 13.4 | 0.4 | 1.1 | 0.8 |
| other Harpacticoida | ind m ⁻³ | 1.8 | 4.6 | 0.8 | 0.5 | 1.3 | 37.9 | 2.1 | 1.6 | 2.2 |
| Monstrilloida | ind m ⁻³ | 0.2 | 0.4 | 0.1 | 0.0 | 0.9 | 2.9 | 0.1 | 0.1 | 0.4 |
| Siphonostomatoida | ind m ⁻³ | 0.0 | 0.6 | 2.2 | 0.0 | 0.2 | 0.2 | 0.0 | 0.0 | 0.1 |
| Copepoda nc | ind m ⁻³ | 2.1 | 1.4 | 0.9 | 0.8 | 0.8 | 2.9 | 3.0 | 0.3 | 0.7 |
| Amphipoda | ind m ⁻³ | 3.5 | 0.1 | 0.1 | 0.0 | 2.2 | 1.3 | 0.1 | 0.6 | 0.2 |
| Chaetognata | ind m ⁻³ | 0.1 | 0.0 | 0.0 | 0.0 | 0.0 | 29.0 | 2.4 | 0.2 | 0.1 |
| Appendicularia | ind m ⁻³ | 229.0 | 50.0 | 29.0 | 0.3 | 14.0 | 85.0 | 2.2 | 1.3 | 127.0 |
| Meroplankton | | | | | | | | | | |
| Proportion of meroplankton | % | 5.6 | 27.3 | 42.2 | 51.1 | 74.1 | 22.4 | 4.4 | 32.8 | 17.0 |
| Gastropoda larvae | ind m ⁻³ | 3.0 | 7.9 | 8.0 | 0.0 | 2.7 | 26.9 | 6.2 | 11.2 | 1.3 |
| Bivalvia | ind m ⁻³ | 2.6 | 2.2 | 0.8 | 0.1 | 2.7 | 11.7 | 0.9 | 0.7 | 2.6 |
| Polichaeta larvae | ind m ⁻³ | 8.8 | 8.2 | 4.4 | 3.6 | 15.0 | 76.9 | 19.8 | 11.5 | 6.0 |
| Platyhelminthes larvae | ind m ⁻³ | 0.0 | 0.3 | 0.9 | 0.1 | 69.9 | 0.7 | 0.0 | 0.2 | 0.0 |
| Nemertea larvae | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| Cirripedia nauplia | ind m ⁻³ | 5.8 | 82.5 | 31.6 | 0.5 | 8.4 | 37.4 | 0.6 | 2.5 | 29.5 |
| Decapoda larvae | ind m ⁻³ | 3.1 | 12.8 | 1.5 | 0.2 | 0.3 | 0.9 | 0.8 | 0.2 | 2.8 |
| Mysidacea | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.8 | 0.0 | 0.0 | 0.0 |
| Cumacea | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.1 | 0.0 | 1.0 | 0.0 | 0.1 | 0.0 |
| Tanaidacea | ind m ⁻³ | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.3 | 0.0 | 0.0 | 0.0 |
| Isopoda | ind m ⁻³ | 0.3 | 0.1 | 0.0 | 0.1 | 0.0 | 0.8 | 0.6 | 0.1 | 0.1 |
| Phoronida | ind m ⁻³ | 0.0 | 0.2 | 0.1 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| Ascidiacea larvae | ind m ⁻³ | 0.4 | 10.9 | 0.6 | 0.3 | 3.9 | 3.7 | 0.4 | 2.1 | 3.3 |
| Echinodermata larvae | ind m ⁻³ | 0.0 | 0.7 | 0.5 | 0.0 | 0.0 | 0.0 | 0.2 | 0.9 | 1.1 |
| Ichthyoplankton | ind m ⁻³ | 1.1 | 1.3 | 0.5 | 4.6 | 0.4 | 1.2 | 11.3 | 4.8 | 5.6 |

Table 3. Results of 2-way-ANOVAs of physical, chemical and biological variables by port and sampling period (Cagliari: C, El Kantaoui: E, Heraklion: H) and 1-way-ANOVA by stations in Cagliari port.

| Variable | 2-way-ANOVA by port and sampling period | | Scheffé mean multicomparison by port main effect | | | Scheffé mean multicomparison by sampling period main effect | | | 1-way-ANOVA | Scheffé mean multicomparison by stations in Cagliari | | | | | |
|---|---|----------------|--|----|----|---|-----|-----|--------------|--|----|----|----|----|----|
| | Significance | r ² | C | E | H | Feb | May | Sep | Significance | r ² | C1 | C2 | C3 | C4 | C5 |
| number of observations | | | 15 | 9 | 14 | 12 | 13 | 13 | | | 3 | 3 | 3 | 3 | 3 |
| ENVIRONMENT | | | | | | | | | | | | | | | |
| Temperature | *** | 0.95 | b | a | a | c | b | a | ns | | | | | | |
| Salinity | *** | 0.59 | b | a | a | | | | ** | 0.76 | b | a | a | a | ab |
| DO | *** | 0.68 | a | b | b | a | a | b | ns | | | | | | |
| pH | ** | 0.37 | | | | b | ab | a | ns | | | | | | |
| Ammonia (NH ₄) | * | 0.26 | a | b | b | | | | ** | 0.80 | a | b | b | b | b |
| Nitrite (NO ₂) | ** | 0.37 | a | b | b | | | | *** | 0.83 | a | b | b | b | b |
| Nitrate (NO ₃) | ** | 0.35 | a | b | ab | | | | ** | 0.73 | a | ab | b | b | ab |
| Phosphate (PO ₄) | * | 0.28 | a | b | b | | | | ** | 0.77 | a | b | b | b | ab |
| Silicate (SiO ₂) | * | 0.32 | a | b | ab | | | | ** | 0.81 | a | b | b | b | ab |
| POC | *** | 0.52 | a | b | b | b | a | ab | ns | | | | | | |
| Chlorophyll- <i>a</i> | *** | 0.54 | a | b | b | b | a | ab | ns | | | | | | |
| AHs | * | 0.26 | a | ab | b | | | | ns | | | | | | |
| PAHs | *** | 0.32 | a | b | b | a | b | b | ** | 0.18 | ab | a | ab | b | ab |
| BACTERIO-, PHYTO- & ZOO-PLANKTON | | | | | | | | | | | | | | | |
| Total bacterial counts | *** | 0.57 | a | b | b | | | | ns | | | | | | |
| Total phytoplankton | *** | 0.43 | a | b | b | | | | ns | | | | | | |
| Diatoms | *** | 0.43 | a | b | b | | | | ns | | | | | | |
| Dinoflagellates | *** | 0.46 | | | | b | a | b | ns | | | | | | |
| Coccolithophores | *** | 0.51 | ab | b | a | a | b | b | ns | | | | | | |
| Other phytoplankton | ** | 0.35 | a | b | b | . | | | ns | | | | | | |
| Total zooplankton | ns | | | | | | | | ns | | | | | | |

r² = coefficient of determination of the ANOVA model. In each of the three multi-comparisons, different letters indicate significant differences at p<0.05. Letter 'a' represents a higher value than letter 'b' that represents a higher value than letter 'c'. Ns and empty spaces: not significant (p>0.05). DO: dissolved oxygen; POC: particulate organic carbon; AHs: aliphatic hydrocarbons; PAHs: polycyclic aromatic hydrocarbons.

Table 4. Results of factor analysis of the physical-chemical data and log-transformed densities of planktonic organisms in three Mediterranean ports. Factor coefficients in bold were used for interpretation. n=190 observations.

| Variable | Coefficients | | | |
|-------------------------------------|---------------------------------------|----------------------------|----------------------------------|-----------------------------|
| | Factor1 | Factor2 | Factor3 | Factor4 |
| ENVIRONMENT | | | | |
| Temperature | -0.01 | 0.34 | -0.77 | 0.12 |
| Salinity | -0.87 | 0.06 | -0.21 | 0.10 |
| DO | 0.58 | -0.18 | 0.26 | 0.25 |
| Ammonia | 0.76 | -0.24 | 0.25 | 0.02 |
| Nitrite | 0.86 | -0.20 | 0.21 | 0.09 |
| Nitrate | 0.65 | -0.11 | 0.49 | 0.02 |
| Phosphate | 0.79 | -0.22 | 0.13 | 0.12 |
| Silicate | 0.68 | -0.14 | 0.50 | 0.12 |
| POC | 0.76 | 0.04 | -0.17 | 0.38 |
| Chlorophyll-a | 0.85 | -0.02 | -0.15 | 0.31 |
| AHs | 0.40 | -0.11 | 0.21 | -0.44 |
| PAHs | 0.16 | -0.19 | 0.36 | -0.42 |
| BACTERIA & PHYTOPLANKTON | | | | |
| Total bacterial counts | 0.70 | 0.20 | 0.04 | 0.01 |
| Diatoms | 0.77 | 0.01 | -0.53 | -0.11 |
| Dinoflagellates | -0.19 | 0.06 | -0.73 | 0.27 |
| Coccolitophores | -0.38 | -0.09 | 0.48 | 0.23 |
| Other phytoplankton | 0.79 | -0.11 | -0.36 | -0.01 |
| ZOOPLANKTON | | | | |
| Holoplankton | | | | |
| Cnidaria Scyphomedusae | 0.04 | 0.18 | 0.32 | -0.28 |
| Cnidaria Siphonophora | -0.35 | -0.01 | 0.41 | 0.49 |
| Ostracoda | 0.04 | 0.18 | 0.03 | 0.08 |
| Amphipoda | 0.03 | 0.05 | -0.04 | -0.47 |
| Cladocera | 0.65 | 0.40 | 0.20 | -0.21 |
| Calanoida (Copepoda) | -0.13 | 0.67 | 0.31 | 0.38 |
| Cyclopoida (Copepoda) | -0.03 | 0.76 | 0.47 | -0.05 |
| Harpacticoida (Copepoda) | 0.07 | 0.85 | 0.00 | -0.13 |
| Monstrilloida (Copepoda) | -0.05 | 0.59 | -0.09 | -0.09 |
| Siphonostomatoida (Copepoda) | 0.43 | -0.17 | -0.30 | 0.15 |
| Chaetognata | -0.24 | 0.65 | 0.17 | -0.04 |
| Appendicularia | 0.46 | 0.58 | 0.09 | -0.26 |
| Meroplankton | | | | |
| Cnidaria Hydromedusae | 0.09 | 0.76 | 0.09 | -0.16 |
| Ascidiacea | 0.36 | 0.35 | -0.37 | 0.18 |
| Platyhelmintha | -0.05 | -0.09 | -0.56 | -0.29 |
| Polychaeta larvae | -0.14 | 0.71 | 0.08 | 0.04 |
| Gasteropoda larvae | 0.09 | 0.58 | -0.03 | 0.33 |
| Bivalvia larvae | 0.02 | 0.74 | -0.04 | -0.19 |
| Echinodermata larvae | 0.00 | 0.03 | -0.12 | 0.43 |
| Cirripedia nauplia | 0.58 | 0.46 | -0.35 | 0.03 |
| Decapoda larvae | 0.46 | 0.41 | 0.00 | 0.28 |
| Ichthyoplankton | -0.44 | 0.07 | 0.33 | 0.52 |
| Variance Explained (%) | 24 | 15 | 11 | 7 |
| Interpretation | Freshwater input, nutrient loading | Zooplankton variability | Seasonality, water turbulence | Anthropogenic activities |

DO: dissolved oxygen; POC: particulate organic carbon; AHs: aliphatic hydrocarbons; PAHs: polycyclic aromatic hydrocarbons. The zooplankton taxa are those groups in Table 2b that presented at least 50 ind m⁻³ in the whole dataset.

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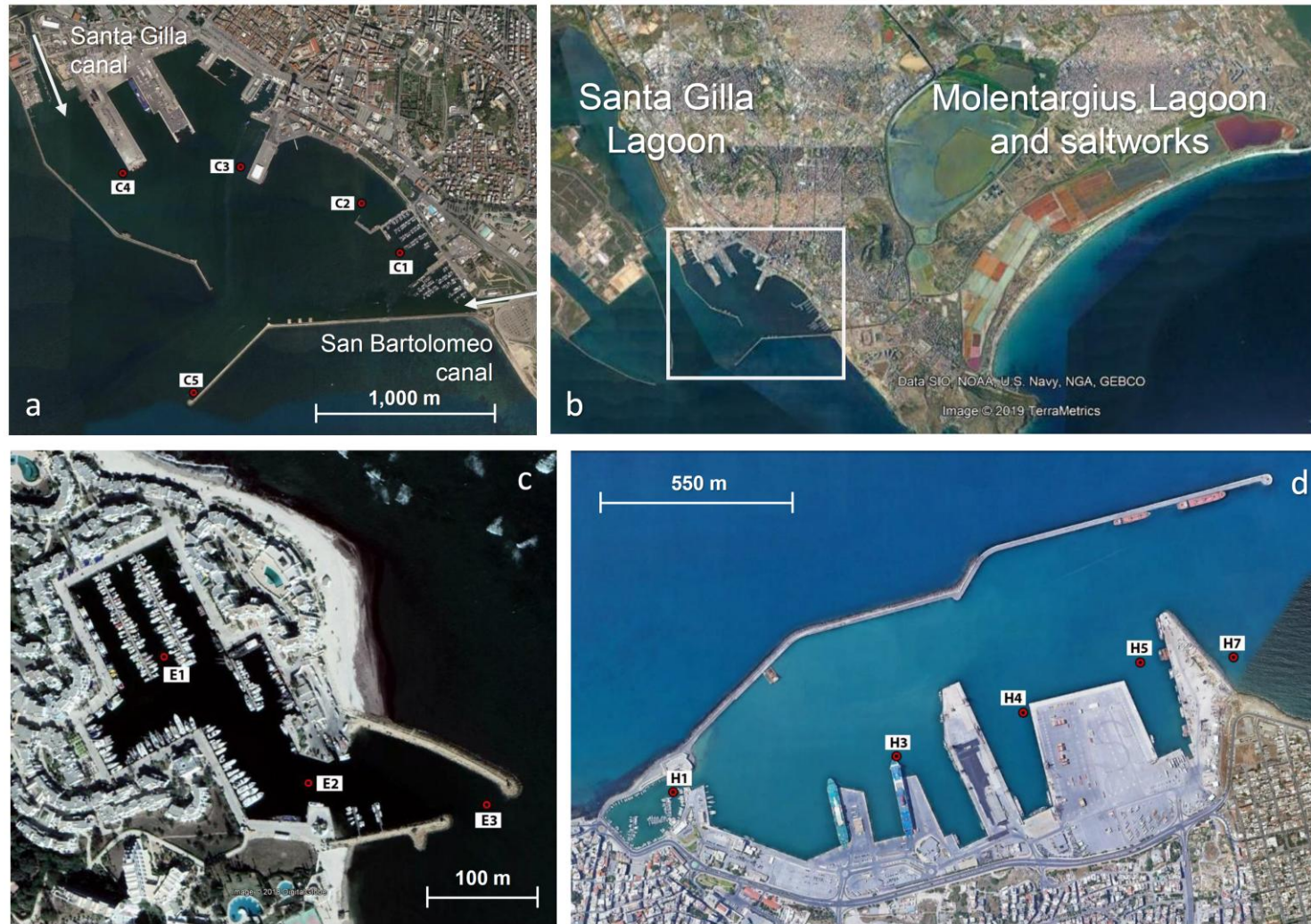
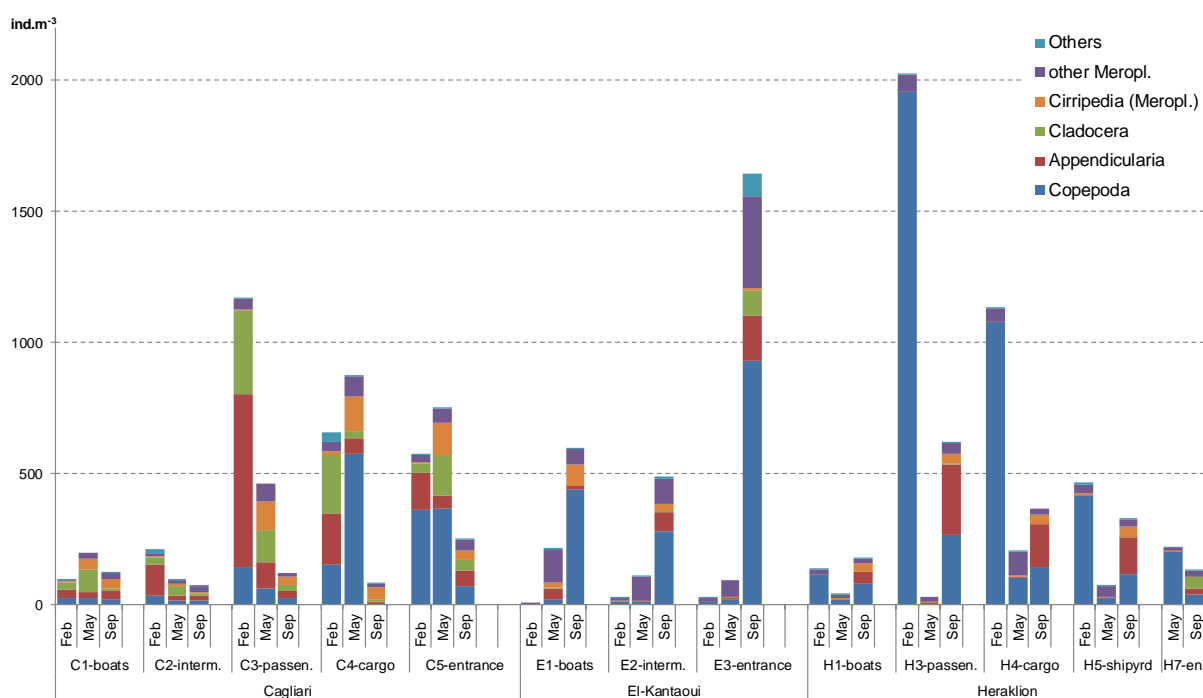
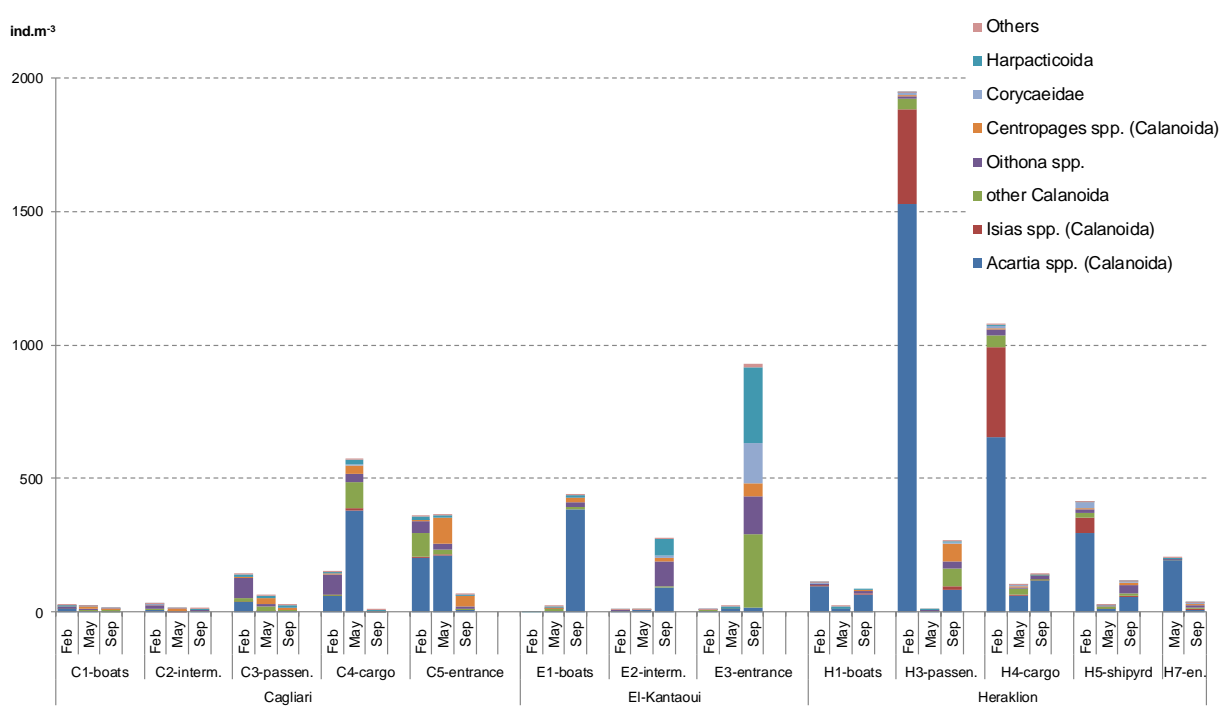


Figure 1. Maps of the three Mediterranean ports and sampling stations. **a)** Port of Cagliari (Italia, Sardinia) and inlets of freshwaters (arrows); **b)** Systems of lagoons and canals surrounding the Port of Cagliari (in the box); **c)** Port of El Kantaoui (Sousse, Tunisia); **d)** Port of Heraklion (Crete, Heraklion) (Google Earth © 2017, v. 7.1.8.3036).

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6 **Figure 2.** Densities of the zooplankton taxonomic groups that mostly contributed to the total in different
1 7 stations and sampling periods. **a)** Total zooplankton. For clarity of representation only the components
2 8 contributing more than 5% to the total zooplankton were represented. (Copepoda 67%; Appendicularia
3 9 12%; Cladocera 7%; Cirripedia 5%; other Meroplankton 7%; others 2%). **b)** Copepods. For clarity of
4 10 representation only the components contributing more than 1% to the copepods were represented.
5 11 (*Acartia* 67%; *Isias* 14%; other Calanoida 6%; *Oithona* 5%; *Centropages* 4%; Corycaeidae 1%; Harpacticoida
6 12 2%; others < 1%) Explanations of the labels and station uses are in Table 1. Further details in text.

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Figure 3. Variation of each factor through space (ports, stations) and time (seasons); the most relevant variables for each factor, based on the coefficients in **Table 4**, are reported on the left. **a)** Factor 1; **b)** Factor 2; **c)** Factor 3; **d)** Factor 4. Arrows indicate the presence of both positive and negative coefficients (Factors 1,3,4) or only positive (Factor 2). Explanations of the labels and station uses are in Table 1.

Supplementary Data 1S

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Supplementary Data 2S

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Declaration of interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

*Credit Author Statement

CR, ET, and FS developed the concept for this study. CR, ET, and FS carried out the field work. ET performed the bacterioplankton analysis. CN performed phytoplankton analysis. CR performed zooplankton analysis. AM and CR performed statistical analyses. All authors interpreted the results, drafted the manuscript, reviewed the final version of the manuscript and approved it before submission.