

SCIENTIA MARINA 83(1)
March 2019, 41-52, Barcelona (Spain)
ISSN-L: 0214-8358
<https://doi.org/10.3989/scimar.04836.16A>

Summer mesozooplankton assemblages in relation to environmental parameters in Kavala Gulf, northern Aegean Sea

Aikaterini Karagianni, Anna Artemiou, Athanassios C. Tsikliras, Evangelia Michaloudi

Department of Zoology, School of Biology, Aristotle University of Thessaloniki, Thessaloniki, Greece.

(AK) E-mail: aikatekm@bio.auth.gr. ORCID iD: <https://orcid.org/0000-0003-3004-8868>

(AA) E-mail: anna.artemiou@gmail.com. ORCID iD: <https://orcid.org/0000-0002-7161-6879>

(ACT) E-mail: atsik@bio.auth.gr. ORCID iD: <https://orcid.org/0000-0002-9074-3259>

(EM) (Corresponding author) E-mail: tholi@bio.auth.gr. ORCID iD: <https://orcid.org/0000-0001-8194-2865>

Summary: Shallow coastal areas are ecosystems with high productivity. Although the eastern part of the Mediterranean Sea is oligotrophic, the shallow coastal waters of the northern Aegean, such as Kavala Gulf, are productive due to the influence of the Black Sea water and the presence of freshwater input from three rivers. The aim of this work was to determine the structure of zooplankton communities in Kavala Gulf in the summer of 2002 and 2003 and to investigate their relation to environmental variables. Zooplankton communities were characterized by the presence of common coastal Cladocera, such as *Penilia avirostris*, small pelagic Copepoda, such as the calanoida *Acartia clausi* and the cyclopoida *Oithona plumifera*, and Tunicata, such as *Oikopleura*, *Fritillaria* and Doliolidae. The abundances corresponded to the peak of the warm period and were significantly greater in 2002 because of a *P. avirostris* bloom, which seemed to have better exploited the environmental sources favouring its dominance in the area. Overall, the structure of summer mesozooplankton communities in Kavala Gulf follows the pattern exhibited by mesozooplankton communities in other Greek coastal areas of the northern Aegean Sea.

Keywords: abiotic factors; quantitative distribution; ecological associations; diversity patterns; coastal zone; zooplankton community composition.

Comunidades zooplanctónicas estivales en relación con parámetros ambientales en el golfo de Kavala, norte del mar Egeo

Resumen: Las aguas costeras poco profundas son ecosistemas con alta productividad. Aunque el mar Mediterráneo oriental es oligotrófico, las aguas costeras poco profundas al norte del mar Egeo, como el golfo de Kavala, son productivas debido a la influencia de aguas provenientes del mar Negro y a los aportes de agua dulce procedentes de tres ríos. El objetivo de este trabajo fue determinar la estructura de las comunidades de zooplancton en el golfo de Kavala durante los veranos de 2002 y 2003, e investigar su relación con variables ambientales. Las comunidades de zooplancton se caracterizaron por la presencia de cladóceros costeros comunes, como *Penilia avirostris*, pequeños copépodos pelágicos, tales como el calanoide *Acartia clausi* y el ciclopoide *Oithona plumifera*, y tunicados como *Oikopleura*, *Fritillaria* y Doliolidae. Las abundancias correspondieron al pico del periodo cálido y fueron significativamente mayores en 2002 debido a una proliferación de *P. avirostris*, que parece que supo explotar mejor las condiciones ambientales favoreciendo su dominio en la zona. En general, la estructura de las comunidades de mesozooplancton de verano en el golfo de Kavala siguen el patrón exhibido por las comunidades de mesozooplancton en otras áreas costeras griegas al norte del mar Egeo.

Palabras clave: factores abióticos; distribución cuantitativa; asociaciones ecológicas; patrones de diversidad; zona costera; composición de la comunidad zooplanctónica.

Citation/Como citar este artículo: Karagianni A., Artemiou A., Tsikliras A.C., Michaloudi E. 2019. Summer mesozooplankton assemblages in relation to environmental parameters in Kavala Gulf, northern Aegean Sea. *Sci. Mar.* 83(1): 41-52. <https://doi.org/10.3989/scimar.04836.16A>

Editor: E. Saiz.

Received: July 26, 2018. **Accepted:** December 14, 2018. **Published:** March 1, 2019.

Copyright: © 2019 CSIC. This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International (CC BY 4.0) License.

INTRODUCTION

Zooplankton play an important role in marine pelagic food webs, connecting primary producers, small pelagic fish populations and benthic communities, and thus transferring organic matter through the pelagic food webs (Fenchel 1988). Plankton and fish larvae communities are often structured in assemblages that are closely related to environmental characteristics (Cowen et al. 1993). Their response to alterations of the environmental conditions can be detected by observing changes in diversity, community structure and species distribution (e.g. Dortch et al. 1989, Richardson 2008). Moreover, zooplankton population dynamics can be indicative of environmental changes and may affect higher trophic levels (Beaugrand et al. 2003, Frederiksen et al. 2006). As an important part of the pelagic food web and fisheries ecology, zooplankton has thus been used in the evaluation and modelling of marine ecosystems (e.g. Triantafyllou et al. 2007, Petihakis et al. 2009).

The northern Aegean Sea, in the northeastern Mediterranean, is a complex and dynamic marine ecosystem (Tsagarakis et al. 2010). Its hydrodynamic complexity is closely linked to the influence of the Black Sea water masses, which outflow in the upper part of the water column through the Dardanelles Straits and are characterized by low salinity and temperature (Zodiatis and Balopoulos 1993, Zervakis and Georgopoulos 2002). Almost permanently, one branch of the Black Sea current follows a northward direction towards the Thracian Sea, where it is captured by an anticyclonic gyre formed around the island of Samothraki and is mainly restricted to the upper 0 to 20 m layer (Zodiatis and Balopoulos 1993, Zervakis and Georgopoulos 2002). The water circulation pattern in the northern Aegean has a considerable influence on the horizontal oceanographic variability of zooplankton assemblages (Isari et al. 2006, 2007, Zervoudaki et al. 2006). These hydrological conditions lead to high productivity, in contrast with the rest of the eastern Mediterranean Sea (Lykousis et al. 2002), making the northern Aegean one of the most productive areas in the region (Bosc et al. 2004).

Kavala Gulf and the surrounding area constitutes one of the most well-known fishing and nursery grounds of the northern Aegean Sea (Stergiou et al. 1997), which is particularly important for small pelagic fish populations such as European anchovy (*Engraulis encrasicolus*), the European pilchard (*Sardina pilchardus*) and the round sardinella (*Sardinella aurita*) (Stergiou et al. 2007, Sylaios et al. 2010), which may compete for resources (Tsikliras 2014). As zooplankton is the basic component in the diet of the early stages of fish and of the adults of small pelagic fish, zooplankton communities in Kavala Gulf are key organisms of the pelagic web (Tsikliras et al. 2005, Catalán et al. 2010, Karachle and Stergiou 2013). Although zooplankton communities have been widely studied in the northern Aegean Sea (Table 1), Kavala Gulf was not included among the study areas. Filling this gap could provide a valuable asset, adding to the theoretical knowledge of ecosystem functioning and fisheries modelling that could be used to refine or expand the ecosystem models that have already been developed for the area (Tsagarakis et al. 2010). The aim of the present study was to fill this knowledge gap regarding mesozooplankton in Kavala Gulf through the analysis of a fine grid of 17 stations during two summer cruises in 2002 and 2003. In particular, our analysis aimed to (i) demonstrate the spatial variability of major zooplankton species/taxonomic groups and (ii) investigate the relation between sea surface abiotic factors and the structure of zooplankton communities.

MATERIALS AND METHODS

Study site

Kavala Gulf (40.88056°N, 24.41667°E) is a semi-enclosed area located on the continental shelf of the northern Aegean Sea in the eastern Mediterranean Sea (Fig. 1). It is connected to the Aegean Sea through its main mouth in the south, which is wide (20 km) and deep, and in the east, through the smaller (7.3 km wide), shallower mouth, the strait of the island of Thassos (Fig. 1). As Kavala Gulf is shallow (mean depth:

Table 1. – List of selected publications on marine mesozooplankton in the Aegean Sea.

Area	Year	Sampling frequency	Net mesh size	Citation
Thermaikos Gulf	1984-1985	Seasonal	200 µm	Siokou-Frangou and Papathanassiou 1991
	1988-1990	Monthly	200 µm	Alvanou 1999
Saronikos Gulf	1984-1985	Monthly	200 µm	Siokou-Frangou 1996, Siokou-Frangou et al. 1998
	1987	Monthly	200 µm	Siokou-Frangou et al. 2009
	1989-1993	Monthly	200 µm	Christou 1998, Christou and Stergiou 1998, Siokou-Frangou et al. 2009
	1998	Monthly	200 µm	Siokou-Frangou et al. 2009
N Aegean	1997	Seasonal	200 µm	Siokou-Frangou et al. 2002, Lykousis et al. 2002
	1998	Seasonal	200 µm	Lykousis et al. 2002
	1999	Snapshot	200 µm	Zervoudaki et al. 2006
	2000	Snapshot	200 µm	Zervoudaki et al. 2006
	2003	Snapshot	250 µm	Isari et al. 2008
	2004	Snapshot	250 µm	Isari et al. 2008
	2005	Snapshot	250 µm	Isari et al. 2008
	2006	Snapshot	250 µm	Isari et al. 2008
	2008	Snapshot	200 µm	Frangoulis et al. 2010
	2009	Snapshot	200 µm	Siokou et al. 2014
2011	Monthly	200 µm	Siokou et al. 2014	
C Aegean	1998	Snapshot	250 µm	Ramfos et al. 2005

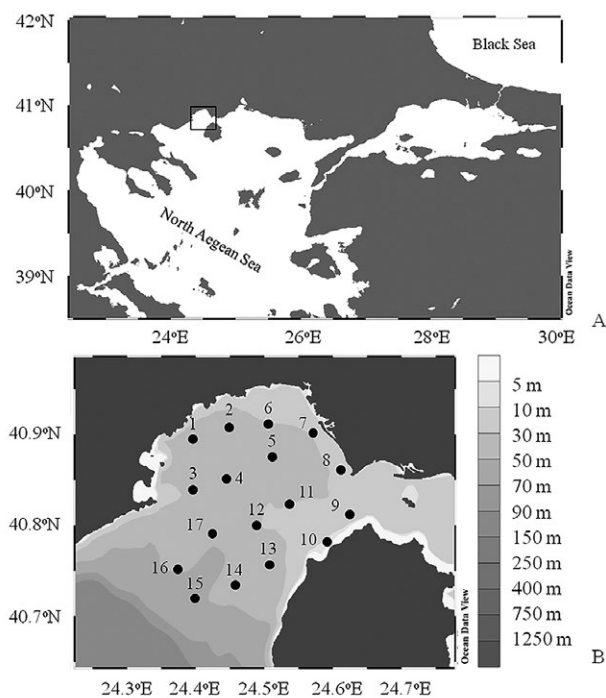


Fig. 1. – A, map of the northern Aegean Sea. Black square indicates Kavala Gulf. B, map of the sampling stations in Kavala Gulf, northern Aegean Sea in July 2002 and 2003.

34 m, maximum depth: 50 m), with a flat seabed, the topographical variations inside it are minimal (Fig. 1). The hydrology of the gulf is influenced by the fresher and cooler Black Sea water, which oscillates seasonally in occurrence, intensity and direction (Sylaios et al. 2005), and forms a surface current flowing to the northern coastline of Greece (Poulos et al. 1997). Kavala Gulf also receives fresh water inflows from the Nestos River, though these have been significantly reduced since the construction of two hydroelectric dams in 1997 (Sylaios et al. 2005).

Sampling process and laboratory analysis

Mesozooplankton samples and environmental data (depth, temperature, salinity and dissolved oxygen) were collected during two sampling periods in Kavala Gulf in the summers of 2002 and 2003. The surveys were carried out across a fine scale grid of 17 stations (Fig. 1) during the period of thermal stratification, which coincides with the spawning period of European anchovy (*Engraulis encrasicolus*) and round sardinella (*Sardinella aurita*). These two small pelagic clupeoids, together with the winter spawning European pilchard (*Sardina pilchardus*), are the main zooplanktivorous fish in the area (Somarakis 2005, Tsikliras et al. 2005). Based on their depth and distance from the shore, the 17 stations had been previously grouped (Tsikliras and Koutrakis 2011) into shallow (maximum depth less than 30 m: stations 6, 7, 8, 9, 10) and deep (maximum depth above 30 m: stations 1, 2, 3, 4, 5, 11, 12, 13, 14, 15, 16, 17).

Sea surface temperature, salinity and dissolved oxygen were measured at each station using CTD sensors (Ocean 301, Idronaut). The hydrographic sam-

pling profiles of all these parameters are available in Tsikliras et al. (2009). For the mesozooplankton collection, a paired bongo net sampler was used, with two 0.6 m diameter frames fitted with 250 μ m mesh conical nets. The cod-end consisted of a plastic container incorporating a window of 250 μ m mesh net that allowed water to escape. A flowmeter was centrally mounted on the opening of each frame to the net to estimate the volume of water (m^3) flowing through the net. The sampler was deployed in a double oblique tow from the surface down to 1 to 2 m from the bottom, thus forming a V-shaped dive profile. Retrieval speed was kept at the same rate (20 $m\ min^{-1}$) for all samplings, which were all carried out in daylight (between 09:00 h and 18:00 h). Upon recovery of the sampler, the plankton was washed into a jar and fixed with 4% buffered formalin solution.

After the removal of fish larvae for further investigation of ichthyoplankton assemblages (Tsikliras et al. 2009, Tsikliras and Koutrakis 2011, Tsikliras 2014), the major mesozooplankton taxa (e.g. Copepoda, Cladocera, Doliolidae, Appendicularia, etc) were identified using the appropriate taxonomic keys. Copepoda (only adults) and Cladocera were further identified down to species level when possible, whereas copepodites were grouped together and were not further identified. For the abundance estimation, each sample was enumerated in aliquots under the stereoscope. Subsamples were taken with a pipette, their size ranging from 1/2 to 1/30 of the total sample depending on the sample's density. The total number of counted organisms was on average 1844.47 ± 646.58 ind.

Data analysis

Mesozooplankton abundance data were expressed as number of individuals per volume unit (m^3) by dividing the individuals per sample by the volume of the filtered water. The standardized numbers were used to calculate the percentage contribution of each taxon to the total abundance. In order to identify the similarity of mesozooplankton communities among stations and years, hierarchical cluster analysis based on the Bray-Curtis similarity index (Bray and Curtis 1957) was performed on the mesozooplankton taxa's abundance matrix. Abundance data were $\log(x+1)$ transformed in order to approach normality and homogeneity of variances and to reduce bias due to highly abundant groups. CLUSTER was run using group-average linking. The similarity profile (SIMPROF) permutation test option (default settings of 999 permutations and significance level 0.05) was applied to indicate significant groups in the resulting dendrogram. The similarity analysis routines, analysis of similarity (ANOSIM) and similarity percentage analysis (SIMPER) were used to test the significance levels and sources of variance between the various zooplankton assemblages associated with the groupings identified in the hierarchical cluster analysis. The above analyses were conducted with the Plymouth Routine In Multivariate Ecological Research (PRIMER) v.5 software package (Clarke and Gorley

2001). Permutational multivariate analysis of variance (PERMANOVA), based on Euclidean distance resemblance matrix, was used to test for statistical differences in the mesozooplankton abundances between the sampled periods (July 2002 and 2003) and between water depth categories (stations at less than 30 m and more than 30 m depth). Analyses were performed using the Paleontological Statistics software package (Past v.3.15) (Hammer et al. 2001).

Direct ordination analyses were used to assess significant relationships between biological and environmental data. All environmental variables (temperature, salinity, depth and dissolved oxygen) were $\log(x+1)$ transformed. Previously, a detrended correspondence analysis (DCA) was performed and, as biological data showed a linear response with respect to environmental gradients, a redundancy analysis (RDA) was applied. All environmental parameters with an inflation factor smaller than 20 were included in the analysis as explanatory variables (Ter Braak and Verdonschot 1995); the abundance of each zooplankton taxon was included as a response variable. The statistical significance of the variation in the parameters and the overall significance of the ordination were tested with the Monte Carlo permutation test (as default settings of 499 unrestricted permutations; $P < 0.05$). Ordination analyses were performed using the CANOCO program, version 4.5 (Ter Braak and Šmilauer 2002).

RESULTS

Environmental parameters

The values of sea surface environmental parameters (temperature, salinity, oxygen) and maximum depth of the sampling stations in Kavala Gulf are given in Table 2. Salinity and oxygen differed significantly between the years (ANOVA; $P < 0.005$), whereas temperature did not (ANOVA; $P > 0.005$).

Community composition and structure

Overall, a total of 9 main holoplanktonic and 6 main meroplanktonic groups were recorded (Table 3). Nineteen taxa were further identified, 12 of them down to species level and 7 to genus level (Table 3).

Total abundance ranged from 41069 to 164939 ind m^{-3} in 2002 and from 5508 to 86747 ind m^{-3} in 2003 (Fig. 2A). Total abundance values were significantly higher in 2002 than in 2003 at all sampling stations (pseudo- $F = 7.27$, $P < 0.05$) (Fig. 2B) except station 10, where abundance was higher in 2003 (Fig. 2A). No significant difference was observed between shallow and deeper stations (pseudo- $F = 0.14$, $P > 0.05$) according to the PERMANOVA procedure. Cladocera were the dominant group at all sampling stations, followed by Copepoda and Appendicularia in rank order, whose cumulative contribution exceeded 79% of mesozooplankton abundance (Table 3). Among the four identified Cladocera species, *Penilia avirostris* showed a marked predominance (Fig. 3A, B), followed by *Evadne nordmanni* Lovén, 1836, *Evadne spinif-*

Table 2. – Sea surface physical-chemical parameters and maximum depth of the 17 stations in Kavala Gulf. Stations' abbreviations are consistent with the formula ST (number of station) (sampling year; 02 for 2002 and 03 for 2003).

Year	Station	Temperature (°C)	Salinity	Oxygen	Depth (m)
2002	ST0102	24.3	34.0	8.2	37.0
	ST0202	24.2	33.8	8.5	35.2
	ST0302	23.7	33.7	7.6	34.0
	ST0402	23.9	33.5	8.4	43.0
	ST0502	24.4	33.9	8.4	37.0
	ST0602	25.1	33.8	8.7	29.0
	ST0702	25.5	33.8	8.6	25.0
	ST0802	24.3	32.9	8.5	28.0
	ST0902	24.5	34.0	8.4	28.0
	ST1002	24.3	34.3	8.5	16.0
	ST1102	24.6	33.2	8.6	30.0
	ST1202	24.5	33.3	7.8	30.0
	ST1302	25.1	34.3	8.0	30.0
	ST1402	25.0	34.2	8.3	35.0
	ST1502	24.6	34.4	8.3	45.0
	ST1602	25.0	34.3	8.0	43.0
	ST1702	25.2	34.2	8.1	44.5
2003	ST0103	24.7	31.6	8.3	36.5
	ST0203	24.8	32.5	8.1	36.0
	ST0303	25.3	31.5	7.9	43.0
	ST0403	25.3	32.4	7.8	40.5
	ST0503	24.4	31.3	8.5	38.5
	ST0603	24.2	32.5	8.3	27.8
	ST0703	24.2	31.4	8.3	25.0
	ST0803	24.9	30.5	7.9	27.0
	ST0903	25.4	32.6	7.8	28.0
	ST1003	25.3	32.7	7.7	13.6
	ST1103	24.6	32.9	8.1	36.5
	ST1203	24.7	32.7	8.0	30.0
	ST1303	25.0	32.7	8.0	36.0
	ST1403	24.7	30.6	8.3	39.0
	ST1503	24.3	30.5	8.4	45.0
	ST1603	24.4	32.4	8.3	45.0
	ST1703	25.4	32.6	7.9	41.0

era P.E. Müller, 1867 and *Pseudevadne tergestina* (Claus, 1877) (Table 3). Despite small differences in the species-specific distribution patterns, the bulk of Cladocera were found more or less in the eastern part of the gulf in 2002 and in the shallow stations of the southeastern part near Thassos Island in 2003 (Fig. 3A, B). Copepods, being the second dominating taxonomic group, were mainly represented by juveniles and small-sized zooplankters [e.g. the calanoida *Acartia (Acartiura) clausi* Giesbrecht, 1889 and *Paracalanus parvus* (Claus, 1863), the cyclopoida *Oithona plumifera* Bair 1843 and *Corycaeus* spp. Dana 1845, and the harpacticoida *Macrosetella* spp. Scott A., 1909] (Table 3), whose distribution did not follow a discrete pattern (Fig. 3C, D). Copepods' mean contribution to mesozooplankton abundance was higher in 2003 than in 2002 (16.07% and 11.10%, respectively) (Table 3). Appendicularia were represented by the genus *Fritillaria* Fol, 1872, and the genus *Oikopleura* Mertens, 1830, which contributed 2.92% and 1.87% in 2002 and 3.01% and 2.20% in 2003, respectively (Table 3). The hierarchical cluster analysis (Fig. 4) revealed that the mesozooplankton community structure differed significantly between stations with varying maximum depth and sampling period. Four groups were generated (ANOSIM, $R = 0.746$, $P = 0.001$; Appendix 1), with no apparent pattern between shallow and deep stations (Fig. 4). Station 10 fell into different groups (Groups A and D in

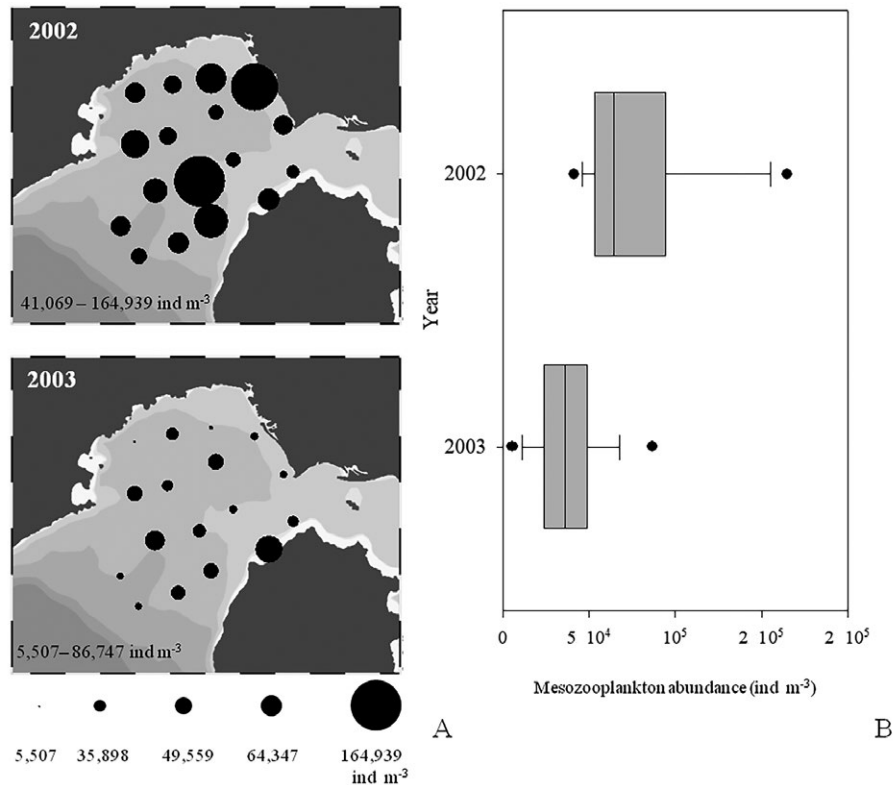


Fig. 2. – A, distribution of mesozooplankton abundance in the water column in Kavala Gulf, northern Aegean in July 2002 and 2003. Values for smaller and larger disc (range of abundance values) are indicated. B, box plots of mesozooplankton abundance in Kavala Gulf, northern Aegean in July 2002 and 2003 grouped by sampling year. Outliers are plotted as dots.

Table 3. – List of recorded taxa. Mean abundance values (Abu) and mean relative abundance % (Abu %) in the summer of 2002 and 2003 in Kavala Gulf. Abb, abbreviation of each taxon as used for further analyses; Ca, Calanoida; Cy, Cyclopoida; Ha, Harpacticoida.

Taxa	Abb	Abu		Abu %	
		2002	2003	2002	2003
Holoplankton					
Amphipoda	Amp	242.99	0.00	0.58	0.00
Appendicularia	App	3155.50	1759.59	4.46	5.12
<i>Fritillaria</i> spp. Fol, 1872	Fri	1979.21	1043.77	2.92	3.01
<i>Oikopleura</i> spp. Mertens, 1830	Oik	1249.80	715.82	1.87	2.20
Chaetognatha	Sag	194.61	155.24	0.38	0.49
Cladocera	Cla	60358.22	26220.55	76.01	68.14
<i>Evadne nordmanni</i> Lovén, 1836	Evn	2517.24	1032.31	3.11	3.26
<i>Evadne spinifera</i> P.E.Müller, 1867	Evs	1073.36	300.84	1.41	1.00
<i>Penilia avirostris</i> (Dana, 1852)	Pea	55344.58	24135.30	47.61	48.58
<i>Pseudevadne tergestina</i> (Claus, 1862)	Pst	1423.04	752.09	1.89	2.29
Copepoda	Cop	8087.28	5130.61	11.10	16.07
<i>Acartia (Acartiura) clausi</i> Giesbrecht, 1889 (Ca)	Acc	1986.01	1659.70	2.62	5.55
<i>Calocalanus pavo</i> (Dana, 1849) (Ca)	Cap	48.31	0.00	0.52	0.00
<i>Centropages typicus</i> Kroeyer, 1849 (Ca)	Cet	1175.98	455.20	1.93	1.60
<i>Conaea rapax</i> Giesbrecht, 1891 (Cy)	Cor	31.98	32.55	0.17	0.44
<i>Corycaeus</i> spp. Dana, 1845 (Cy)	Crc	265.92	200.21	0.33	0.65
<i>Ctenocalanus vanus</i> (Dana, 1849) (Ca)	Ctv	737.62	455.60	1.08	1.58
<i>Macrosetella</i> spp. Scott A., 1909. (Ha)	Mac	0.00	8.38	0.00	0.51
<i>Mecynocera</i> spp. Thompson I.C., 1888 (Ca)	Mec	206.53	81.17	0.47	0.54
<i>Oithona plumifera</i> Bair, 1843 (Cy)	Oip	1192.12	1010.17	1.74	2.93
<i>Paracalanus parvus</i> (Claus, 1863) (Ca)	Pap	750.34	222.76	1.14	0.63
<i>Temora stylifera</i> (Dana, 1848) (Ca)	Tes	550.51	235.25	0.83	0.67
unidentified Copepoda	Cop	1472.83	924.01	2.10	2.89
Doliolidae	Dol	3063.61	917.41	4.17	2.35
Ostracoda	Ost	355.36	23.78	0.52	0.80
Medusae	Hyd	611.48	380.50	0.90	1.14
Siphonophores	Sip	731.62	2156.19	1.12	6.00
Meroplankton					
Decapoda larvae	Dec	269.15	126.34	0.60	0.42
Echinoderma	Ech	455.16	94.48	0.74	0.43
Gastropoda	Gas	64.68	0.00	0.44	0
Mysidae larvae	Mys	2625.70	0.00	3.77	0.00
Polychaeta	Pol	51.21	62.67	0.61	0.40
Cirripedia	Cir	169.28	133.76	0.25	0.43
Total		77610.34	37060.20		

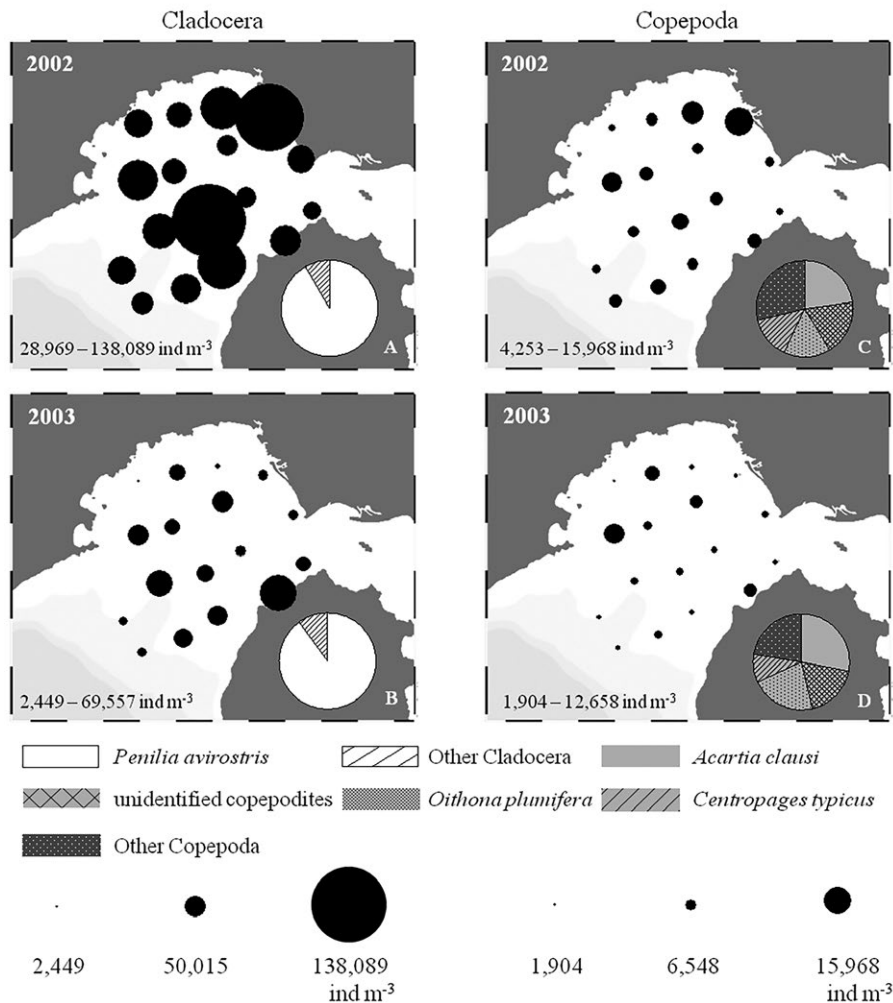


Fig. 3. – Abundance of Cladocera (A and B) and Copepoda (C and D) community in Kavala Gulf during 2002 (A and C) and 2003 (B and D). Discs are proportional to abundance. Values for smaller and larger disc (range of abundance values) are indicated. In the right corner of each graph the average between the sampling stations' contribution of dominant taxa in the community of Cladocera (A and C) and Copepoda (B and D) is plotted in a pie graph.

2002 and 2003, respectively) (Fig. 4). This was mainly due to Mysidae, Ostracoda and Amphipoda abundance for Group A (SIMPER, average dissimilarity $\leq 27.18\%$; Appendix 1) and to Cirripedia, Echinoderma, Decapoda and Polychaeta abundance for Group D (SIMPER, average dissimilarity 14.48% ; Appendix 1). However, according to ANOSIM, these groupings had no statistical significance (ANOSIM, Group A, $R \geq 0.986$, $P > 0.05$ and Group D, $R = 0.645$, $P > 0.05$; Appendix 1). The rest of the sampling stations of 2002 and station 9 of 2003 were grouped together in the upper group of the cluster (Group B), while the stations of 2003 were placed in Group C (Fig. 4). Groups B and C differed significantly, as shown by pairwise comparison (ANOSIM, $R = 0.703$, $P = 0.01$; Appendix 1). Their dissimilarity was mainly due to the contribution of Ostracoda, Echinoderma, Polychaeta and Doliolidae (SIMPER, average dissimilarity 15.06% ; Appendix 1), whereas the similarity among stations in the same groups was in both cases mainly due to Cladocera, Copepoda, Appendicularia and Doliolidae or Siphonophora in rank order (SIMPER, average similarity Group B 91.50% and Group C 89.72% ; Appendix 1).

Relationships between physical-chemical parameters and zooplankton community structure

To assess significant relationships between surface environmental data and mesozooplankton community structure, ordination analysis of taxa assemblages, expressed in terms of biomass, was conducted (Fig. 5). In the diagnostic DCA, the highest value of the length of gradient of axis was 0.819, indicating that the relationship between mesozooplankton and environmental variables was linear (Ter Braak and Šmilauer 2002), and an RDA was performed. The significant environmental variables ($P < 0.05$) included in the RDA were salinity and maximum depth. Although dissolved oxygen was not a significant variable ($P = 0.06$), it was included in the analysis as an explanatory variable of zooplankton distribution. The Monte Carlo test confirmed that the selected RDA model was significant (F ratio = 9.531, $P = 0.002$). The eigenvalues of the first two axes were 0.241 and 0.072, and both of them together explained 90.4% of the variation in species-environment relation. The first axis, which accounted for a total variance of 69.6% , was negatively and strongly correlated with

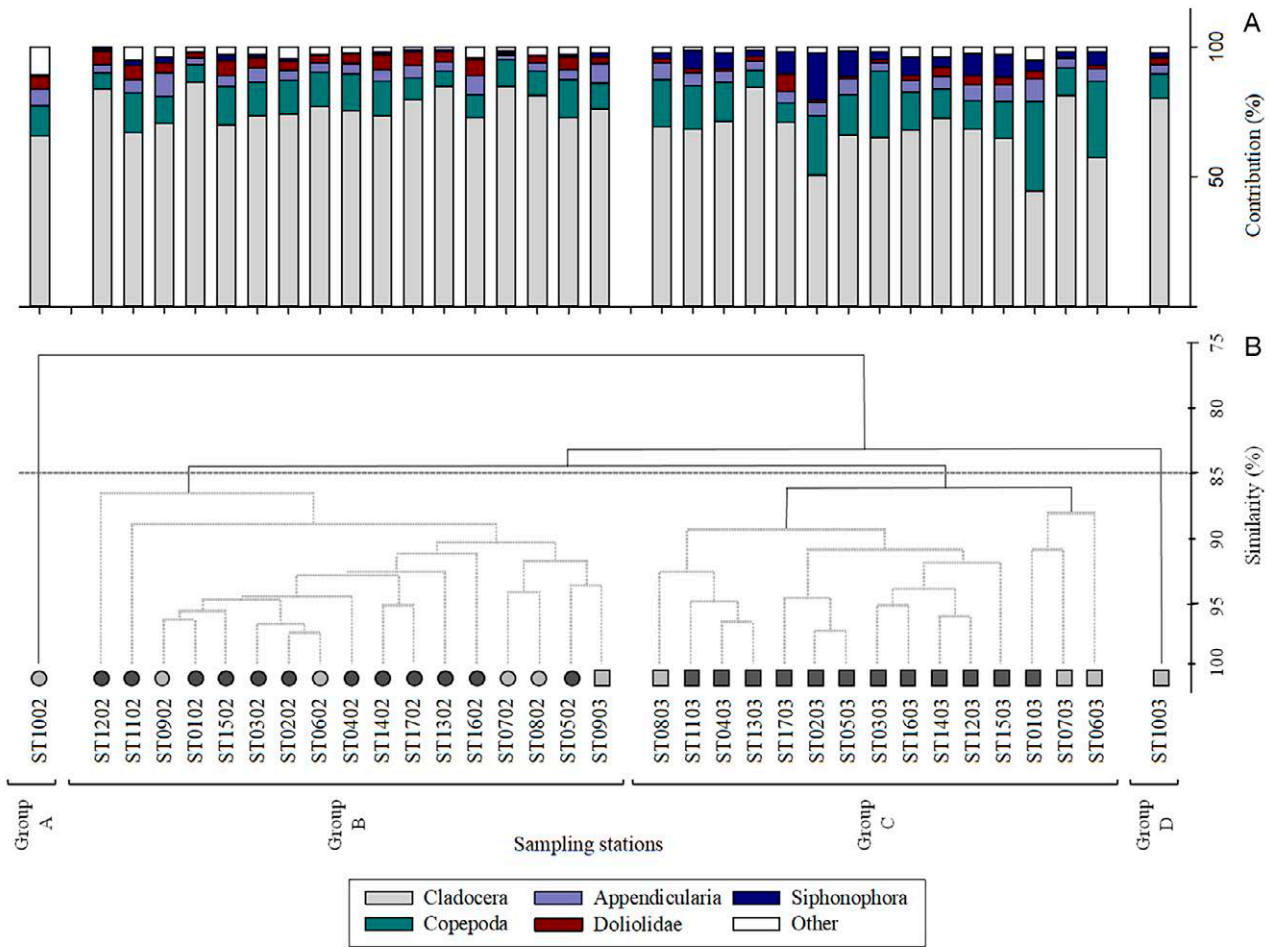
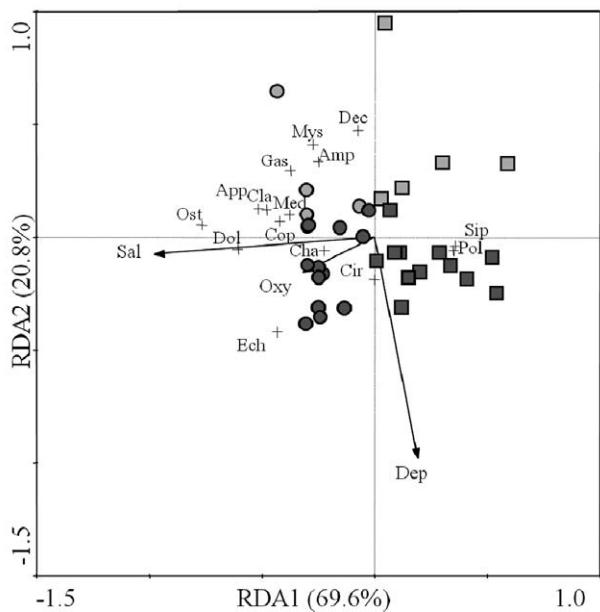


Fig. 4. – A, percentage contribution of dominant mesozooplankton groups to total abundance in a grid of 17 stations in Kavala Gulf in summer 2002 and 2003. B, dendrogram of Bray-Curtis similarity index of mesozooplankton community among samplings in sampling grid in Kavala Gulf from hierarchical agglomerative cluster analysis in log x+1 transformed data of zooplankton group abundance. Solid black lines indicate significant branches based on SIMPROF permutation tests, while dashed light grey lines indicate non-significant divisions (P<0.05). Symbols coded for sampling year and maximum depth categories; light grey circles for shallow stations in 2002; light grey squares for shallow stations in 2003 and dark grey squares for deeper stations in 2003. Sampling abbreviations are consistent with the formula ST (number of station) (sampling year; 02 for 2002 and 03 for 2003).



sea surface salinity ($r=0.9768$). Moreover, mesozooplankton communities in 2002, being in contrast with those of 2003, were negatively correlated with Axis 1 (Appendix 2) and consequently positively correlated with sea surface salinity. Axis 2 showed 20.8% variation and was strongly and negatively correlated with maximum depth ($r=-0.9768$). Dissolved oxygen was strongly and negatively correlated with the third axis ($r=-0.9355$), which accounted for a total of 9.6% variation. Cumulative fit indicated that the predominant mesozooplankton groups affiliated with Ostracoda and

Fig. 5. – Triplot diagram of mesozooplankton major taxonomic groups; explanatory variables and samples in a redundancy analysis (RDA) of the first and second axis illustrating the relationships among communities in the grid of stations in Kavala Gulf in the summer of 2002 and 2003. Square and circle points indicate the position of mesozooplankton communities in relation to environmental parameters (solid arrows) and species composition (cross points) based on taxon abundance. Mesozooplankton communities from each sampling were coded based on year of sampling and maximum depth same as in Figure 4B. Taxa abbreviations are shown in Table 3.

Cladocera displayed stronger correlations with Axis 1 than the rest of the groups (Appendix 3). Ostracoda, Doliolidae, Cladocera, Appendicularia and Mysidae were notable taxonomic groups with higher correlations with Axis 2 (Appendix 3).

DISCUSSION

The present study focuses on the community structure of mesozooplankton in Kavala Gulf in the summer of 2002 and 2003. Although our dataset is relatively old, this is to the best of our knowledge the first published study on mesozooplankton in Kavala Gulf. Mesozooplankton sampling was part of a fisheries survey in the area, whose data on hydrology and fisheries have already been published (Tsikliras et al. 2005, 2009, Tsikliras 2014). Since Kavala Gulf is an important fisheries and nursery ground (Stergiou et al. 1997), data on mesozooplankton could contribute to the theoretical knowledge of ecosystem functioning in the area. Moreover, though our dataset lacks seasonality, it could be useful for comparisons with present and future data, and for further meta-analyses in fisheries and ecosystem modelling through the refining or expanding of models that have already been developed for the area (Tsagarakis et al. 2010).

The hydrological data collected during the present study were limited to the surface layer. Nevertheless, they are in accordance with other studies in the northern Aegean (e.g. Isari et al. 2008). The differences recorded between the two sampling years for salinity are attributed to the rainier days just before the 2003 survey (Tsikliras et al. 2009). The composition of the zooplankton community found in the present study is typical of coastal areas in the Aegean and Ionian Seas, such as Saronikos Gulf (Siokou-Frangou 1996), Patraikos Gulf, the northern Euboikos Gulf, Pagasitikos Gulf (Ramfos et al. 2005) and the Thracian Sea (Isari et al. 2007). Moreover, a similar zooplankton community composition has been recorded during the summer in other coastal ecosystems of the Mediterranean and Black Seas (Razouls et al. 2018). As sampling was originally planned for ichthyoplankton collection, zooplankton samples were not obtained with the typical WP-2 net but with a net of a wider mesh size (250 µm). Thus, differences in community composition are expected to be mainly in small-sized species, due to selectivity of large-sized species by the gear. Furthermore, 250 mesh size nets have also been used for studies in the northern Aegean Sea (Table 1), making our results still comparable. Seasonal fluctuations in zooplankton abundance of the coastal Mediterranean regions are characterized by two maxima, one in spring and one in autumn (Stergiou et al. 1997). In the Mediterranean coastal areas, the spring peak is common in the seasonal fluctuations of zooplankton abundance (Siokou-Frangou 1996), while the autumn peak is observed mainly in areas strongly affected by anthropogenic activities, often during the period June-September (Siokou-Frangou 1996, Alvanou 1999). We acknowledge that our dataset is limited to the summer period and is not able to reflect the annual fluctuations

of mesozooplankton communities. Nevertheless, the mesozooplankton abundance values recorded, after comparison with summer data in other coastal areas of the Mediterranean Sea, seemed to correspond to the autumn peak (Kovalev et al. 2003). In order to broaden our knowledge on Kavala Gulf a more intensive annual sampling scheme should be designed in the future.

Total mesozooplankton abundance values in both sampling years were relatively higher than those of samplings in the surrounding area of the Thracian Sea (Isari et al. 2006) and in other coastal systems in the Mediterranean [e.g. Gulf of Naples (Mazzocchi and Ribera d'Alcalà 1995); Gulf of Trieste (Lipej et al. 1997); Ionian and central Aegean Sea (Ramfos et al. 2005)] and the Black Sea (e.g. Sazhina 1964). This may be linked to the special conditions occurring in this semi-enclosed gulf, namely the effects of the Black Sea water, the nutrient rich intake from the nearby river outflows and the urban and industrial activity in the area (Friligos 1985, Kardaras 2005, Sylaios et al. 2005, Isari et al. 2006). A water circulation study that was synchronous with our sampling in Kavala Gulf states that northern Aegean water enters westwards in the Gulf from the Strait of Thassos (near station 10, Fig. 1), forming a cyclonic sea surface pattern (Tsikliras et al. 2009). Moreover, in Kavala Gulf cold water masses with low salinity are provided by the outflow of the Nestos River during winter and spring (Kardaras 1998), whereas in the summer period water circulation is influenced by the low-salinity water masses entering from the Dardanelles (Kardaras 1998, Tsikliras et al. 2009). The influence of the Black Sea water on zooplankton communities has been previously studied in the northern Aegean Sea, showing that it favours the dominance of Cladocera and small-sized Copepoda due to its lower salinity (Isari et al. 2006, 2007, 2011), as is the case with the mesozooplankton communities of Kavala Gulf. Moreover, the abundance recorded in 2002 was higher than in 2003, while the reverse pattern is observed in the ichthyoplankton data (Tsikliras et al. 2009). This reverse relationship may be indicative of fish larvae predation on the mesozooplankton communities (Cushing 1990). However, inter-annual variability of the mesozooplankton-ichthyoplankton trophic link and longer time-series would be needed to support this hypothesis with more evidence.

The dominant zooplankton group in both sampling years were Cladocera, followed by Copepoda and Tunicata (Fig. 4). Cladocera dominate coastal areas in the summer period according to similar studies, contributing to the summer peak [e.g. Christou and Stergiou 1998; Strymonikos and Ierissos Gulfs (Michaloudi 1999); Thracian Sea (Isari et al. 2007)]. They are favoured by high temperatures, low salinity (Moraitou-Apostolopoulou and Kiortsis 1973, Christou and Stergiou 1998) and low depth (Siokou-Frangou 1996), which are conditions encountered in Kavala Gulf during the summer. This trend was reflected in the RDA (Fig. 5), with stations being grouped by depth and surface salinity. In 2002, mesozooplankton communities were positively correlated with the sea surface salinity values, which were significantly higher, whereas in 2003

they were negatively correlated. Although salinity was significantly differentiated between the years studied, it was not an actual forming factor of the communities' distinctness because of the narrow range of the measured values (1.89 ± 0.95). According to Siokou-Frangou et al. (1998), temperature is the main driving factor of mesozooplankton community formation.

In our study, the Cladocera community consisted almost entirely of *P. avirostris*, which is a highly efficient filter feeder feeding mainly on phytoplankton, such as diatoms, nanoflagellates (HNF) and bacterioplankton (Turner et al. 1988, Kim et al. 1989, Atienza et al. 2006). *P. avirostris* peaks may be favoured by the presence of *Cymodocea* and *Posidonia* meadows in Kavala Gulf (Orfanidis et al. 2010), taking advantage of the dissolved organic carbon enrichment in the pelagic food web and the microbial loop that is favoured in such habitats (Barrón et al. 2004, Barrón and Duarte 2009). Moreover, massive populations of *P. avirostris* may be a result of the combination of its ability to exploit the available food resources and of its life cycle characteristics (a short life cycle and parthenogenetic reproduction) (Colton 1985, Valentin and Marazzo 2003). In conclusion, the zooplankton community of a semi-enclosed gulf in the northern Aegean Sea during the summer was mainly dominated by Cladocera, small pelagic Copepoda and Tunicata. The high abundance values might have been favoured by the Black Sea water and river input in the upper layer of the water column. Our results on the mesozooplankton communities of Kavala Gulf may provide valuable information concerning the function of the pelagic food web in the gulf and contribute to the construction and/or improvement of models used in fisheries management of economically important fish stocks of the area.

ACKNOWLEDGEMENTS

We thank N. Kamidis, A. Argyrokastritis and P. Karanikolas for field work at sea, and D. Dimarchopoulou for proofreading the manuscript. The present work is part of a research project (PENED 2001) funded by the General Directorate of Research and Technology, Ministry of Development (Greece).

REFERENCES

- Alvanou L. 1999. Annual cycle of Cladocerans in the Thermaikos Gulf and the Gulf of Thessaloniki. *Fresenius Environ. Bull.* 8: 724-731.
- Atienza D., Calbet A., Saiz E., et al. 2006. Trophic impact, metabolism, and biogeochemical role of the marine cladoceran *Penilia avirostris* and the co-dominant copepod *Oithona nana* in NW Mediterranean coastal waters. *Mar. Biol.* 150: 221-235. <https://doi.org/10.1007/s00227-006-0351-z>
- Barrón C., Duarte C.M. 2009. Dissolved organic matter release in a *Posidonia oceanica* meadow. *Mar. Ecol. Prog. Ser.* 374: 75-84. <https://doi.org/10.3354/meps07715>
- Barrón C., Marbé N., Terrados J., et al. 2004. Community metabolism and carbon budget along a gradient of seagrass (*Cymodocea nodosa*) colonization. *Limnol. Oceanogr.* 49: 1642-1651. <https://doi.org/10.4319/lo.2004.49.5.1642>
- Baugrand G., Brander K.M., Lindley J.A., et al. 2003. Plankton effect on cod recruitment in the North Sea. *Nature* 426: 661-664. <https://doi.org/10.1038/nature02164>
- Bosc E., Bricaud A., Antoine D. 2004. Seasonal and interannual variability in algal biomass and primary production in the Mediterranean Sea, as derived from 4 years of SeaWiFS observations. *Global Biogeochem. Cycles* 18. <https://doi.org/10.1029/2003GB002034>
- Bray J.R., Curtis J.T. 1957. An ordination of the upland forest communities of southern Wisconsin. *Ecol. Monogr.* 27: 325-349. <https://doi.org/10.2307/1942268>
- Catalán I.A., Folkvord A., Palomera I., et al. 2010. Growth and feeding patterns of European anchovy (*Engraulis encrasicolus*) early life stages in the Aegean Sea (NE Mediterranean). *Est. Coast. Shelf Sci.* 86: 299-312. <https://doi.org/10.1016/j.ecss.2009.11.033>
- Christou E.D. 1998. Interannual variability of copepods in a Mediterranean coastal area (Saronikos Gulf, Aegean Sea). *J. Mar. Syst.* 15: 523-532. [https://doi.org/10.1016/S0924-7963\(97\)00080-8](https://doi.org/10.1016/S0924-7963(97)00080-8)
- Christou E.D., Stergiou K.I. 1998. Modelling and forecasting the fortnightly cladoceran abundance in the Saronikos Gulf (Aegean Sea). *J. Plankton Res.* 20: 1313-1320. <https://doi.org/10.1093/plankt/20.7.1313>
- Clarke K.R., Gorley R.N. 2001. *PRIMER v5: User Manual/ Tutorial*. Primer-E Ltd., Plymouth, 91 pp.
- Colton Jr. J.B. 1985. Eastward extension of the distribution of the marine cladoceran (*Penilia avirostris*) in the northwest Atlantic: a case of ecesis. *J. Northwest Atl. Fish. Soc.* 6: 141-148. <https://doi.org/10.2960/J.v6.a14>
- Cowen R.K., Hare J.A., Fahay M.P. 1993. Beyond hydrography: can physical processes explain larval fish assemblages within the Middle Atlantic Bight? *Bull. Mar. Sci.* 53: 567-587.
- Cushing D.H. 1990. Plankton production and year-class strength in fish populations: an update of the match/mismatch hypothesis. *Adv. Mar. Biol.* 26: 249-293. [https://doi.org/10.1016/S0065-2881\(08\)60202-3](https://doi.org/10.1016/S0065-2881(08)60202-3)
- Dortch Q., Packard T.T. 1989. Differences in biomass structure between oligotrophic and eutrophic marine ecosystems. *Deep Sea Res. Part 1. Oceanogr. Res. Pap.* 36: 223-240.
- Fenchel T. 1988. Marine plankton food chains. *Annu. Rev. Ecol. Syst.* 19: 19-38. <https://doi.org/10.1146/annurev.es.19.110188.000315>
- Frangoulis C., Psarra S., Zervakis V., et al. 2010. Connecting export fluxes to plankton food-web efficiency in the Black Sea waters inflowing into the Mediterranean Sea. *J. Plankton Res.* 32: 1203-1216. <https://doi.org/10.1093/plankt/fbq010>
- Frederiksen M., Edwards M., Richardson A.J., et al. 2006. From plankton to top predators: bottom-up control of a marine food web across four trophic levels. *J. Anim. Ecol.* 75: 1259-1268. <https://doi.org/10.1111/j.1365-2656.2006.01148.x>
- Friligos N. 1985. Nutrient conditions in two Aegean gulfs. *Mar. Pollut. Bull.* 16: 208-209. [https://doi.org/10.1016/0025-326X\(85\)90482-5](https://doi.org/10.1016/0025-326X(85)90482-5)
- Hammer Ø., Harper D.A.T., Ryan P.D. 2001. *PAST: Paleontological statistics software package for education and data analysis*. *Palaeontol. Electron.* 4.
- Isari S., Ramfos A., Somarakis S., et al. 2006. Mesozooplankton distribution in relation to hydrology of the Northeastern Aegean Sea, Eastern Mediterranean. *J. Plankton Res.* 28: 241-255. <https://doi.org/10.1093/plankt/fbi058>
- Isari S., Psarra S., Pitta P., et al. 2007. Differential patterns of mesozooplankton distribution in relation to physical and biological variables of the northeastern Aegean Sea (eastern Mediterranean). *Mar. Biol.* 151: 1035-1050. <https://doi.org/10.1007/s00227-006-0542-7>
- Isari S., Fragopoulou N., Somarakis S. 2008. Interannual variability in horizontal patterns of larval fish assemblages in the northeastern Aegean Sea (eastern Mediterranean) during early summer. *Estuar. Coast. Shelf Sci.* 79: 607-619. <https://doi.org/10.1016/j.ecss.2008.06.001>
- Isari S., Somarakis S., Christou E.D., et al. 2011. Summer mesozooplankton assemblages in the north-eastern Aegean Sea: the influence of Black Sea water and an associated anticyclonic eddy. *J. Mar. Biol. Ass. U.K.* 91: 51-63. <https://doi.org/10.1017/S0025315410000123>
- Karachle P.K., Stergiou K.I. 2013. Feeding and ecomorphology of three clupeoids in the N Aegean Sea. *Medit. Mar. Sci.* 15: 9-26. <https://doi.org/10.12681/mms.350>
- Kardaras T. 1998. The dynamical response of the water masses due to the influence of the prevailing mesoscale meteorological factors. The case of Kavala Bay (North Aegean). PhD thesis. University of Thessaloniki.

- Kardaras T. 2005. The water circulation in Kavala Bay (North Aegean). *Medit. Mar. Sci.* 6: 5-16.
<https://doi.org/10.12681/mms.182>
- Kim S.W., Onbé T., Yoon Y.H. 1989. Feeding habits of marine cladocerans in the Inland Sea of Japan. *Mar. Biol.* 100: 313-318.
<https://doi.org/10.1007/BF00391145>
- Kovalev A.V., Mazzocchi M.G., Kideys A.E., et al. 2003. Seasonal Changes in the Composition and Abundance of Zooplankton in the Seas of the Mediterranean Basin. *Turk. J. Zool.* 27: 205-219.
- Lipej L., Mozetič P., Turk V., et al. 1997. The trophic role of the marine cladoceran *Penilia avirostris* in the Gulf of Trieste. *Hydrobiologia* 360: 197-203.
<https://doi.org/10.1023/A:1003180030116>
- Lykousis V., Chronis G., Tselepidis A., et al. 2002. Major outputs of the recent multidisciplinary biogeochemical researches undertaken in the Aegean Sea. *J. Mar. Syst.* 34: 313-334.
[https://doi.org/10.1016/S0924-7963\(02\)00064-7](https://doi.org/10.1016/S0924-7963(02)00064-7)
- Mazzocchi M.G., Ribera d'Alcalá M.R. 1995. Recurrent patterns in zooplankton structure and succession in a variable coastal environment. *ICES J. Mar. Sci.* 52: 679-691.
[https://doi.org/10.1016/1054-3139\(95\)80081-6](https://doi.org/10.1016/1054-3139(95)80081-6)
- Michaloudi E. 1999. Mesozooplankton in the Strymonikos and Ierissos Gulfs. In: Koutrakis E., Lazaridou E. (eds). Description of the coastal zone of Strymonikos and Ierissos Gulfs. NA-GREF -Fisheries Research Institute - Greek Biotope/Wetland Centre, Nea Peramos, Kavala. [In Greek, English abstract.], pp. 441-453.
- Moraitou-Apostolopoulou M., Kiortsis V. 1973. The cladocerans of the Aegean Sea: occurrence and seasonal variation. *Mar. Biol.* 20: 137-143.
<https://doi.org/10.1007/BF00351451>
- Orfanidis S., Papathanasiou V., Gounaris S., et al. 2010. Size distribution approaches for monitoring and conservation of coastal *Cymodocea* habitats. *Aquat. Conserv.: Mar. Freshw. Ecosyst.* 20: 177-188.
<https://doi.org/10.1002/aqc.1069>
- Petihakis G., Triantafyllou G., Tsiaras K., et al. 2009. Eastern Mediterranean biogeochemical flux model—Simulations of the pelagic ecosystem. *Ocean Sci.* 5: 29-46.
<https://doi.org/10.5194/os-5-29-2009>
- Poulos S.E., Drakopoulos P.G., Collins M.B. 1997. Seasonal variability of the sea surface oceanographic conditions in the Aegean Sea (eastern Mediterranean): an overview. *J. Mar. Syst.* 13: 225-244.
[https://doi.org/10.1016/S0924-7963\(96\)00113-3](https://doi.org/10.1016/S0924-7963(96)00113-3)
- Ramfos A., Somarakis S., Koutsikopoulos C., et al. 2005. Summer mesozooplankton distribution in coastal waters of central Greece (eastern Mediterranean). I. Hydrology and group composition. *J. Mar. Biol. Ass. U.K.* 85: 755-764.
<https://doi.org/10.1017/S0025315405011665>
- Razouls C., de Bovée F., Kouwenberg J., et al. 2005-2018. Diversity and Geographic Distribution of Marine Planktonic Copepods. Accessed 27 June 2018. Available at <http://copepodes.obs-banyuls.fr/en>
- Richardson A.J. 2008. In hot water: zooplankton and climate change. *ICES J. Mar. Sci.* 65: 279-295.
<https://doi.org/10.1093/icesjms/fsn028>
- Sazhina L.I. 1964. Seasonal changes of zooplankton in the north-western part of the Black Sea in 1957. *Proceed. Sevastopol Biol. Stat.* 17: 262-275 (in Russian).
- Siokou-Frangou I. 1996. Zooplankton annual cycle in a Mediterranean coastal area. *J. Plankton Res.* 18: 203-223.
<https://doi.org/10.1093/plankt/18.2.203>
- Siokou-Frangou I., Papathanassiou E. 1991. Differentiation of zooplankton populations in a polluted area. *Mar. Ecol. Prog. Ser.* 76: 41-51.
<https://doi.org/10.3354/meps076041>
- Siokou-Frangou L., Papathanassiou E., Lepretre A., et al. 1998. Zooplankton assemblages and influence of environmental parameters on them in a Mediterranean coastal area. *J. Plankton Res.* 20: 847-870.
<https://doi.org/10.1093/plankt/20.5.847>
- Siokou-Frangou I., Bianchi M., Christaki U., et al. 2002. Carbon flow in the planktonic food web along a gradient of oligotrophy in the Aegean Sea (Mediterranean Sea). *J. Mar. Syst.* 33: 335-353.
[https://doi.org/10.1016/S0924-7963\(02\)00065-9](https://doi.org/10.1016/S0924-7963(02)00065-9)
- Siokou-Frangou I., Zervoudaki S., Christou E.D., et al. 2009. Variability of mesozooplankton spatial distribution in the North Aegean Sea, as influenced by the Black Sea waters outflow. *J. Mar. Syst.* 78: 557-575.
<https://doi.org/10.1016/j.jmarsys.2008.12.025>
- Siokou I., Frangoulis C., Grigoratou M., et al. 2014. Zooplankton community dynamics in the N. Aegean front (E. Mediterranean) in the winter spring period. *Medit. Mar. Sci.* 15: 706-720.
<https://doi.org/10.12681/mms.915>
- Somarakis S. 2005. Marked interannual differences in reproductive parameters and daily egg production of anchovy in the northern Aegean Sea. *Belg. J. Zool.* 134 (Suppl. I): 123-132.
- Stergiou K.I., Christou E.D., Georgopoulos D., et al. 1997. The Hellenic Seas: physics, chemistry, biology and fisheries. *Oceanogr. Mar. Biol. Annu. Rev.* 35: 415-538.
- Stergiou K.I., Moutopoulos D.K., Tsikliras A.C. 2007. Spatial and temporal variability in Hellenic marine fisheries landings. In: Papaconstantinou C., Zenetos A., et al. (eds). State of the Hellenic marine fisheries. Hellenic Centre for Marine Research, Athens, pp. 132-140.
- Sylaos G., Stamatis N., Kallianiotis A., et al. 2005. Monitoring water quality and assessment of land-based nutrient loadings and cycling in Kavala Gulf. *Water Resour. Manag.* 19: 713-735.
<https://doi.org/10.1007/s11269-005-3020-y>
- Sylaos G.K., Koutroumanidis T., Tsikliras A.C. 2010. Ranking and classification of fishing areas using fuzzy models and techniques. *Fish. Manag. Ecol.* 17: 240-253.
<https://doi.org/10.1111/j.1365-2400.2009.00714.x>
- Ter Braak C.J., Šmilauer P. 2002. CANOCO reference Manual and CanoDraw for Windows user's guide: Software for Canonical Community Ordination (version 4.5). Microcomputer Power, Ithaca, New York.
- Ter Braak C.J., Verdonschot P.F. 1995. Canonical correspondence analysis and related multivariate methods in aquatic ecology. *Aquat. Sci.* 57: 255-289.
<https://doi.org/10.1007/BF00877430>
- Triantafyllou G., Korres G., Hoteit I., et al. 2007. Assimilation of ocean colour data into a Biogeochemical Flux Model of the Eastern Mediterranean Sea. *Ocean Sci.* 3: 397-410.
<https://doi.org/10.5194/os-3-397-2007>
- Tsagarakis K., Coll M., Giannoulaki M., et al. 2010. Food-web traits of the North Aegean Sea ecosystem (Eastern Mediterranean) and comparison with other Mediterranean ecosystems. *Est. Coast. Shelf Sci.* 88: 233-248.
<https://doi.org/10.1016/j.ecss.2010.04.007>
- Tsikliras A.C. 2014. Sympatric clupeoid fish larvae in the eastern Mediterranean: coexistence or avoidance? *Adv. Ecol. ID274180*.
<https://doi.org/10.1155/2014/274180>
- Tsikliras A.C., Koutrakis E.T. 2011. Summer fish larval assemblages and station groups in the northern Aegean Sea. *Acta Adriat.* 52: 57-66.
- Tsikliras A.C., Torre M., Stergiou K.I. 2005. Feeding habits and trophic level of round sardinella (*Sardinella aurita*) in the northeastern Mediterranean (Aegean Sea, Greece). *J. Biol. Res.* 3: 67-75.
- Tsikliras A.C., Koutrakis E.T., Sylaos G.K., et al. 2009. Summer distribution of fish larvae in northern Aegean Sea. *J. Mar. Biol. Ass. U.K.* 89: 1137-1146.
<https://doi.org/10.1017/S0025315409000356>
- Turner J.T., Tester P.A., Ferguson R.L. 1988. The marine cladoceran *Penilia avirostris* and the "microbial loop" of pelagic food webs. *Limnol Oceanogr.* 33: 245-255.
<https://doi.org/10.4319/lo.1988.33.2.0245>
- Valentin J.L., Marazzo A. 2003. Modelling the population dynamics of *Penilia avirostris* (Branchiopoda, Ctenopoda) in a tropical bay. *Acta Oecol.* 24: S369-S376.
[https://doi.org/10.1016/S1146-609X\(03\)00021-3](https://doi.org/10.1016/S1146-609X(03)00021-3)
- Zervakis V., Georgopoulos D. 2002. Hydrology and circulation in the north Aegean (eastern Mediterranean) throughout 1997-1998. *Medit. Mar. Sci.* 3: 7-21.
<https://doi.org/10.12681/mms.254>
- Zervoudaki S., Nielsen T.G., Christou E.D., et al. 2006. Zooplankton distribution and diversity in a frontal area of the Aegean Sea. *Mar. Biol. Res.* 2: 149-168.
<https://doi.org/10.1080/17451000600702037>
- Zodiatis G., Balopoulos E. 1993. Structure and characteristics of fronts in the north Aegean Sea. *Boll. Oceanol. Teor. Appl.* 11: 113-124.

APPENDICES

Appendix 1. – Results of the similarity percentage analysis and the pairwise analysis of similarity of variance among the four groups indicated by the hierarchical cluster analysis (Fig. 4B). Taxa with a cumulative contribution of more than 50% are indicated in bold. Taxa abbreviations are shown in Table 3.

	Group A		Group B		Group C		Group D				
	Fewer than 2 samples in group		Contrib (%) Cum (%) Average dissimilarity: 20.29 % (Global R=0.986, P=0.056)		Contrib (%) Cum (%) Average dissimilarity: 27.18 % (Global R=1, P=0.063)		Contrib (%) Cum (%) Average dissimilarity: 22.82 %				
Group A			Mys	22.92	22.92	Mys	18.43	18.43	Mys	21.97	21.97
			Ech	16.72	39.64	Ost	16.06	34.49	Ost	19.73	41.7
			Amp	16.13	55.77	Amp	14.29	48.77	Amp	17.04	58.74
			Gas	13.62	69.4	Gas	12.08	60.85	Cir	15.73	74.47
			Dec	8.53	77.92	Dec	7.61	68.46	Gas	14.4	88.87
			Ost	5.07	82.99	Ech	6.24	74.7	Sip	3.22	92.09
			Cir	3.87	86.87	Dol	4.47	79.17			
			Cha	2.73	89.59	Pol	4.42	83.59			
			Pol	2.58	92.18	Cha	3.21	86.8			
						Sip	2.66	89.47			
					App	2.64	92.11				
Group B			Contrib (%) Cum (%) Average Similarity: 91.50 %		Contrib (%) Cum (%) Average dissimilarity: 15.06 % (Global R=0.703, P=0.01)		Contrib (%) Cum (%) Average dissimilarity: 17.24 % (Global R=0.974, P=0.056)				
			Cla	15.41	15.41	Ost	24.63	24.63	Ech	24.16	24.16
			Cop	12.60	28.01	Ech	15.38	40.01	Ost	22.4	46.56
			App	11.28	39.29	Pol	9.56	49.57	Cir	18.29	64.85
			Dol	10.90	50.19	Dol	8.07	57.64	Dec	8.22	73.06
			Sip	8.93	59.12	Cir	5.83	63.46	Cha	4.61	77.68
			Med	8.52	67.64	Dec	5.6	69.06	Sip	4.32	81.99
			Ech	7.59	75.23	Sip	5.49	74.55	Pol	3.73	85.72
			Ost	7.01	82.24	Cla	5.06	79.61	Med	3.17	88.90
			Cha	6.64	88.88	Cha	4.79	84.4	Amp	2.35	91.24
			Dec	5.61	94.49	Med	3.8	88.2			
						App	3.7	91.89			
	Group C			Contrib (%) Cum (%) Average Similarity: 89.72 %		Contrib (%) Cum (%) Average Dissimilarity: 14.48 % (Global R=0.646, P=0.063)					
			Cla	16.73	16.73	Cir	23.95	23.95			
			Cop	14.37	31.1	Ech	13.63	37.58			
			App	12.29	43.39	Dec	11.7	49.28			
			Sip	11.97	55.36	Pol	9.68	58.96			
			Dol	10.13	65.49	Cha	7.81	66.77			
			Med	9.42	74.9	Dol	7.68	74.45			
			Cir	7.48	82.39	Cla	6.72	81.17			
			Cha	7.07	89.45	Med	6.27	87.44			
			Dec	6.79	96.24	App	4.22	91.66			
Group D	Less than 2 samples in group										

Appendix 2. – Sample scores which are linear combinations of environmental variables.

	Sampling Station	Axis 1	Axis 2	Axis 3	Axis 4
1	ST0102	-0.8411	-0.5768	0.3497	0.0000
2	ST0202	-0.9135	-0.4763	-0.7983	0.0000
3	ST0302	-0.1902	0.0178	25.070	0.0000
4	ST0402	-0.4984	-11.283	-0.2767	0.0000
5	ST0502	-0.9059	-0.6397	-0.4054	0.0000
6	ST0602	-10.944	0.1859	-14.292	0.0000
7	ST0702	-11.123	0.7544	-12.995	0.0000
8	ST0802	-0.2537	0.5127	-10.020	0.0000
9	ST0902	-11.055	0.3737	-0.5790	0.0000
10	ST1002	-15.789	23.624	-0.8948	0.0000
11	ST1102	-0.5673	0.1631	-13.868	0.0000
12	ST1202	-0.0967	0.4432	14.408	0.0000
13	ST1302	-10.722	0.2064	0.7571	0.0000
14	ST1402	-10.931	-0.4368	-0.0021	0.0000
15	ST1502	-11.154	-13.828	0.3497	0.0000
16	ST1602	-0.9109	-11.220	10.375	0.0000
17	ST1702	-0.8897	-12.741	0.6717	0.0000
18	ST0103	10.388	-0.2370	-0.8713	0.0000
19	ST0203	0.4060	-0.2388	0.1755	0.0000
20	ST0303	14.875	-0.6614	0.8881	0.0000
21	ST0403	0.7248	-0.5517	13.264	0.0000
22	ST0503	12.157	-0.4431	-14.167	0.0000
23	ST0603	0.1046	0.6280	-0.7349	0.0000
24	ST0703	10.942	12.159	-0.8678	0.0000
25	ST0803	21.468	11.963	0.1212	0.0000
26	ST0903	0.4371	0.8049	13.587	0.0000
27	ST1003	0.1616	34.563	14.138	0.0000
28	ST1103	0.0236	-0.3742	0.0612	0.0000
29	ST1203	0.2437	0.4425	0.4255	0.0000
30	ST1303	0.3189	-0.2361	0.4735	0.0000
31	ST1403	18.935	-0.3218	-0.8472	0.0000
32	ST1503	19.689	-0.8876	-13.035	0.0000
33	ST1603	0.4317	-11.246	-0.3315	0.0000
34	ST1703	0.5417	-0.6505	10.894	0.0000

Appendix 3. – Cumulative fit per taxon as fraction of variance of species in the redundancy analysis. Taxa with higher correlation (0.25<) with Axis 1 and 2 are marked in bold.

Taxa	Ax1	Ax2	Ax3	Ax4	Var (y)	% Expl
Cladocera	0.2669	0.2830	0.3658	0.5322	0.29	36.58
Copepoda	0.1782	0.1833	0.1981	0.3547	0.12	19.81
Appendicularia	0.2311	0.2461	0.3727	0.5473	0.15	37.27
Chaetognatha	0.0505	0.0537	0.1364	0.1415	0.38	13.64
Siphonophora	0.1264	0.1277	0.1623	0.1925	0.36	16.23
Medusae	0.1423	0.1527	0.1888	0.2908	0.21	18.88
Decapoda	0.0055	0.2294	0.2446	0.3334	0.78	24.46
Cirripedia	0.0000	0.0345	0.1562	0.1562	1.11	15.62
Annelida	0.1219	0.1251	0.1967	0.1985	1.74	19.67
Echinoderma	0.1877	0.3616	0.3664	0.8367	2.87	36.64
Ostracoda	0.5890	0.5922	0.5951	0.8444	3.72	59.51
Dolliolidae	0.3691	0.3716	0.4695	0.6237	0.55	46.95
Mysidacea	0.0755	0.2447	0.2689	0.3017	0.87	26.89
Amphipoda	0.0628	0.1758	0.1870	0.2031	1.14	18.70
Gasteropoda	0.1397	0.2272	0.2508	0.2509	0.73	25.08