



Setting-up a billboard of marine invasive species in the ESENIAS area: current situation and future expectancies

Paraskevi K Karachle, Maria Corsini Foka, Fabio Crocetta, Jakov Dulčić, Nina Dzhembekova, Marika Galanidi, Petya Ivanova, Noa Shenkar, Marius Skolka, Elitsa Stefanova, et al.

► To cite this version:

Paraskevi K Karachle, Maria Corsini Foka, Fabio Crocetta, Jakov Dulčić, Nina Dzhembekova, et al.. Setting-up a billboard of marine invasive species in the ESENIAS area: current situation and future expectancies. *Acta Adriatica*, 2017, 58 (3), pp.429-458. 10.32582/aa.58.3.4 . hal-01976054

HAL Id: hal-01976054

<https://hal.science/hal-01976054>

Submitted on 9 Jan 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Setting-up a billboard of marine invasive species in the ESENIAS area: current situation and future expectancies

Paraskevi K. KARACHLE^{1,*}, Maria CORSINI FOKA², Fabio CROCETTA¹, Jakov DULČIĆ³, Nina DZHEMBEKOVA⁴, Marika GALANIDI⁵, Petya IVANOVA⁴, Noa SHENKAR⁶, Marius SKOLKA⁷, Elitsa STEFANOVA⁴, Kremena STEFANOVA⁴, Victor SURUGIU⁸, Irfan UYSAL⁹, Marc VERLAQUE¹⁰ and Argyro ZENETOS¹

¹ *Institute of Marine Biological Resources and Inland Waters, Hellenic Centre for Marine Research, 46.7 km Athens Sounio ave., P.O. Box 712, 19013 Anavyssos Attiki, Greece*

² *Institute of Oceanography, Hydrobiological Station of Rhodes, Hellenic Center for Marine Research, Cos Street, 85100 Rhodes, Greece*

³ *Institute of Oceanography and Fisheries, Šetalište I. Meštrovića 63, 21000 Split, Croatia*

⁴ *Institute of Oceanology, Bulgarian Academy of Sciences, First May Street 40, P.O.Box 152, Varna 9000, Bulgaria*

⁵ *Institute of Marine Sciences and Technology, Dokuz Eylül University, Haydar Aliyev Bul., No:100, 35430, Inciraltı-Izmir, Turkey*

⁶ *School of Zoology, George S. Wise Faculty of Life Sciences, Tel-Aviv University, Tel-Aviv 6997801, Israel*

⁷ *Ovidius University of Constanta, Bd. Mamaia 124 RO 900527, Constanta, Romania*

⁸ *Alexandru Ioan Cuza University of Iași, Faculty of Biology, Bd. Carol I, No. 20A, 700507 Iași, Romania*

⁹ *Ministry of Forestry & Water Affairs, Marine Protected Areas Division, Alparslan Türkeş cad. No: 72 Bestepe, Ankara, Turkey*

¹⁰ *Aix-Marseille University, Mediterranean Institute of Oceanography (MIO), CNRS/INSU, IRD, UM 110, Campus universitaire de Luminy, case 901, 13288 Marseille cedex 09, France*

Of the 160 species comprising the list, 149 were already present in the ESENIAS countries, while eleven were invasive species either present in the Mediterranean or in other European Seas, likely to be recorded in the ESENIAS countries. The majority of the species were of Red Sea/Indo-Pacific origin (97 species; 60.6%). Italy, Turkey and Greece were the countries with the highest representation of species (159, 152 and 139 species respectively), due to their extended coastline and the number of scholars working on marine invasive species. The highest number of established species was recorded in Turkey (116 species), whereas in Italy and Greece the most numerous species were the “expected” ones (85 and 48 species, respectively). The eastern Adriatic Sea countries (i.e. Albania, Croatia, Montenegro and Slovenia) had generally low numbers of species in this list, many of which are still “expected” to arrive from the neighbouring countries of Greece and Italy. Finally, the most frequently potential pathway was transfer stowaways (ship ballast water: 41 cases; ship hull fouling: 55), whereas unaided spread of Lessepsian immigrants followed (95 cases).

This list is intended to serve as an early warning system that through horizon scanning process would assist ESENIAS countries to prioritise invasive alien species, their pathways and the areas of higher likelihood to appear, in order to take management measures.

Key words: invasive species, ESENIAS, Mediterranean Sea, Black Sea

INTRODUCTION

Biological invasions, both in land and sea, have been worldwide acknowledged as a man-induced ecosystem pressure that should be monitored and managed (ROY *et al.*, 2015, 2017). The Mediterranean, has been characterized as a “sea under siege” (GALIL, 2000), due to the large number of marine alien species recorded, which are introduced by a variety of pathways/vectors (ZENETOS *et al.*, 2010, 2012). In particular, the Eastern Mediterranean, due to its proximity to the Suez Canal, has been susceptible to biological invasions and hosts more than 775 alien and cryptogenic species (ZENETOS *et al.*, 2012). Similarly, the Black Sea, an enclosed marine system of particular physicochemical characteristics, low diversity and high marine traffic, is vulnerable to marine invasions (LEPPÄKOSKI & MIHNEA, 1996). Of particular importance among aliens are the invasive alien species (IAS), which are considered one of the greatest threats to biodiversity and on ecosystem services (KATSANEVAKIS *et al.*, 2014).

Several international agreements such as the Convention on Biological Diversity (CBD), the EU Biodiversity Strategy (EU, 2011), and the European Marine Strategy Framework Directive (EU, 2008), recognise the negative impacts of IAS and highlight the growing concerns of pol-

icy-makers, scientists, stakeholders and society.

It is widely acknowledged among scientists that an early warning system should be elaborated, and one of the best ways is through horizon scanning (ESSL *et al.*, 2015; ROY *et al.*, 2015, 2017; and references therein). Horizon scanning is the process of gathering, analysing and disseminating added-value information to support decision-making (ROY *et al.*, 2014). However, for most regional seas, lists of current or future biological invasions are not available to date. This is also the case for many countries in the Mediterranean region regarding invasive species in the marine environment. Moreover, it is essential to prioritise IAS, their pathways and the areas of higher likelihood to appear, in order to manage and successfully encounter IAS issues (MCGEOCH *et al.*, 2016).

In the present study, effort has been made to compile a list of marine IAS in the network of the ESENIAS (East and South European Network for Invasive Alien Species) countries, including those species already present in this area and those with a high likelihood of appearance. The aim of this work was to create a catalogue (list of invasive / potential invasive species of ESENIAS concern) to serve as a basis for an early warning system, through a horizon scanning process (ROY *et al.*, 2015). This would allow ESENIAS countries to a) identify the most

harmful species within the ESENIAS area; b) support the establishment of an effective early warning and rapid response mechanisms; c) support any eradication attempts of prioritised species at an early stage of invasion, and d) take management measures for IAS that are already widely spread, as requested by the EU Regulation No 1143/2014 (EU, 2014). Detailed fact sheets of the ten most important species, based on their invasive potential, were also assembled.

MATERIAL AND METHODS

In the framework of ESENIAS-Tools project, a list of invasive/potential invasive species in the ESENIAS countries with marine borders (Albania, Bulgaria, Croatia, Greece, Italy, Montenegro, Romania, Slovenia, and Turkey) was built up. The criteria used for this compilation were the following: a) invasive species already present in the area as reported in the national lists or other literature; b) invasive species in the Mediterranean/Black Sea that have not yet exhibited invasive behaviour in ESENIAS countries and c) invasive species in European Seas, which are likely to invade ESENIAS countries. Cryptogenic species were not considered.

Consequently, based on the existing literature, as well as on expert judgment, the species were classified for the ESENIAS area in the countries as established (species with an already established population in the area), casual (species with just one or few records in the country), invasive (species that are established/invasive, with effects – positive and/or negative – on the ecosystem and its services), unknown (species whose presence/status in a country is not yet clarified), and expected (species likely to appear in a country, based on expert judgment and their presence in adjacent areas, and their ecological/environmental requirements). This latter category was further expanded for Croatia and Italy, as expected in the southern and northern parts of the country. The list also includes the species' origin and potential pathway/vector of introduction (CBD, 2014).

Finally, fact sheets were compiled for ten species that were commonly agreed, among

experts, as of high importance. The decision was based on their expanding/invasive character according to biological traits and existing data on their invasion history, combined with the absence of published fact sheets at the time (January 2016), excepting *Mya arenaria* that has recently invaded the Adriatic Sea (CROCETTA & TUROLLA, 2011). Each fact sheet includes the following information: a) description and diagnostic features; b) biology and ecology; c) habitat and distribution (both native and in the ESENIAS area); d) pathway/vector of introduction; e) impacts on biodiversity, ecosystem services and human health; and f) risk assessment and management (when applicable). Maps presented here include information until August 2017. The full fact sheets are published at the ESENIAS Scientific Reports (TRICHKOVA *et al.*, 2017), whereas here only a brief account per species is given.

RESULTS AND DISCUSSION

The list of invasive/potential invasive species

Overall, 160 species are given in the list, 149 of which are already present in the ESENIAS countries and eleven have not yet been recorded in the area. Approximately two thirds of the species (97 species; 60.6%) were of Red Sea/Indo-Pacific origin, followed by those of Pacific origin (28 species; 17.5%) (Table A, on-line supplement). From the species included in this list, the countries with the highest representation were Italy, Turkey and Greece (159, 152 and 139 species respectively; Table 1; Table A, on-line supplement; Fig. 1). A plausible explanation could be the fact that these countries have the longest coastlines among the ESENIAS ones, as well as the highest number of marine experts working on invasive species (KARACHLE *et al.*, 2017). Moreover, a recent work on marinas across the Mediterranean has added considerably to the distribution of many invasive species in Turkey, Greece and Italy (ULMAN *et al.*, 2017).

Turkey was the country with the highest number of established species (116 species), a fact that it can be attributed to its geographic

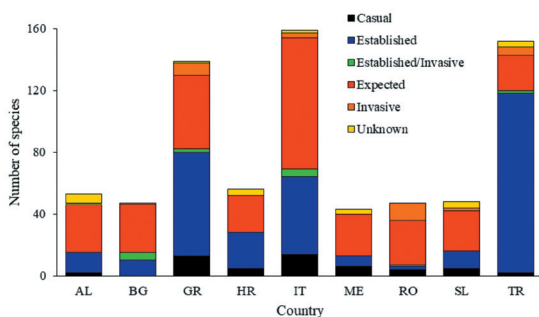


Fig. 1. Number of species per different category in the ESE-NIAS countries with marine borders (AL=Albania, BG=Bulgaria, GR=Greece, HR=Croatia, IT=Italy, ME=Montenegro, RO=Romania, SL=Slovenia, TR=Turkey)

position and proximity to the Suez canal, and, along with Croatia, had the lowest number of expected ones (23 and 24 species, respectively; Table 1; Table A, on-line supplement; Fig. 1). On the other hand, countries with the most numerous expected species were Italy and Greece (85 and 48 species, respectively) followed by Albania, Bulgaria and Romania (31, 31 and 29 species, respectively). In the case of Italy, the high number could be attributed to three main reasons: a) this country is in the middle of the Mediterranean Sea, and hence has influxes of invasive species from both east (mainly Lessepsian immigrants) and west (Atlantic entries); b) intensive oyster activities, mainly in the northern Adriatic, led to accidental transfer of IAS through mariculture, and is thus vulnerable to similar introductions in the future; and c) the country holds major ports and is susceptible to ship transferred species. The last reason stands also for Bulgaria and Romania in the Black Sea. Concerning Greece, it is strongly affected by the Lessepsian invasion, and, due to the proximity to Turkey that holds a substantial number of established alien species, is likely to receive quite many additional IAS.

Finally, a small number of species represented in the list characterizes the eastern Adriatic Sea countries, i.e. Albania, Croatia, Montenegro and Slovenia, in general. Yet, all the aforementioned counties display a high number of “expected” IAS, compared to their total IAS. This fact could be related to the low number

of experts working on alien species in these countries (KARACHLE *et al.*, 2017). In addition, the neighbouring Greece and Italy display high numbers of alien species (e.g. ZENETOS *et al.*, 2015; MARCHINI *et al.*, 2013; ZENETOS & KARACHLE, 2017), many of which have not been reported to date in the East Adriatic countries (KATSANEVAKIS *et al.*, 2011).

The vast majority of the listed species (112 species) had only one potential pathway/vector of introduction, while the remaining ones had either two (42 species) or even three (6 species) pathways/vectors (Table A, on-line supplement). In 94 cases species were transferred as stowaways (ship ballast water: 39 cases; ship hull fouling: 55), whereas in 95 cases species were Lessepsian immigrants.

There were only 15 species transported as contaminants on animals (accidentally with aquaculture) and 10 species by other vectors (aquaria intentional releases, aquaculture escapees) (Table A, on-line supplement). It should be noted that invasive cold water species introduced as contaminants in Northern Europe are likely to be introduced to the Black Sea countries and Northern Adriatic but not to the southern countries. In this context, tropical species that are already invasive in the Levantine are more likely to spread in the southern countries but not in the Black Sea.

Species’ factsheets: a brief account

Ten species were selected, based on their spreading and invasive character, namely *Alexandrium monilatum* (J.F.Howell), *Bonnemaisonia hamifera* Hariot, *Streblospio gynobranchiata* Rice & Levin, 1998, *Cassiopea andromeda* (Forsskål, 1775) upside down jellyfish, *Oithona davisae* Ferrari & Orsi, 1984, *Penaeus aztecus* Ives, 1891 brown shrimp, *Mya arenaria* Linnaeus, 1758 sand gaper, *Diadema setosum* (Leske, 1778) long-spined sea urchin, *Microcosmus exasperatus* Heller, 1878 and *Pterois miles* (Bennett, 1828) Devil firefish/lionfish. The major pathway/vector of introduction for the above mentioned species was transfer stowaways (ship ballast water: 9 cases; ship hull foul-

Table 1. List of present (absent species are indicated by an asterisk *), and expected invasive species in the ESENIAS countries with marine borders (AL=Albania, BG=Bulgaria, GR=Greece, HR=Croatia, IT=Italy, ME=Montenegro, RO=Romania, SL=Slovenia, TR=Turkey). Est=established, cas=casual, unk=unknown establishment status, inv=invasive, exp=expected (expS=expected south; expN=expected north). Empty cells indicate that the species has not been reported yet or is expected in the country. Species in bold are those with factsheets compiled within the ESENIAS-Tools project

Species	AL	BG	GR	HR	IT	ME	RO	SL	TR
<i>Alexandrium monilatum</i> (J.F.Howell)		est		exp	expN		exp		
<i>Caulerpa cylindracea</i> Sonder	est		est	est	inv	est		exp	est
<i>Caulerpa taxifolia</i> (M.Vahl) C.Agardh			exp	est	est			exp	est
<i>Caulerpa taxifolia</i> var <i>distichophylla</i> (Sonder) Verlaque, Huisman and Procacini			est		est				est
<i>Codium fragile</i> (Suringar) Hariot	exp	exp	inv	est	est	exp	exp	est	est
<i>Codium parvulum</i> (Bory de Saint Vincent ex Audouin) P.C.Silva			exp	expS					est
<i>Acrothamnion preissii</i> (Sonder) E.M.Wollaston		exp		est	est		exp	exp	
<i>Asparagopsis armata</i> Harvey	exp		est	est	est	unk		est	est
<i>Asparagopsis taxiformis</i> (Delile) Trevisan de Saint-Léon	est		est	cas	est	est		exp	est
<i>Bonnemaisonia hamifera</i> Hariot	exp	exp	cas	exp	est	exp	exp	cas	est
* <i>Galaxaura rugosa</i> (J.Ellis & Solander) J.V.Lamouroux			exp	expS					unk
<i>Gracilaria vermiculophylla</i> (Ohmi) Papenfuss		exp		exp	est		exp	exp	exp
<i>Lophocladia lallemandii</i> (Montagne) F.Schmitz	est		est	est	est	exp		unk	est
* <i>Polyopes lanceifolius</i> (Harvey) Kawaguchi & Wang		exp		expN			exp		
<i>Womersleyella setacea</i> (Hollenberg) R.E.Norris	exp		est	est	est	est		exp	exp
<i>Sargassum muticum</i> (Yendo) Fensholt		exp			est		exp		
<i>Stypopodium schimperi</i> (Kützinger) M.Verlaque & Boudouresque			est	expS					est
<i>Undaria pinnatifida</i> (Harvey) Suringar		exp	exp	est	est		exp		exp
<i>Halophila stipulacea</i> (Forsskål) Ascherson	est		est	est	est	exp			est
<i>Branchionomma bairdi</i> (McIntosh, 1885)			est		est				est
<i>Branchionomma luctuosum</i> (Grube, 1870)			est		est				est
<i>Ceratonereis mirabilis</i> Kinberg, 1865			exp	expS					est
<i>Desdemona ornata</i> Banse, 1957						exp	exp	exp	est
<i>Dipolydora quadrilobata</i> (Jacobi, 1883)	exp	exp	cas	exp	est			exp	est
<i>Dorvillea similis</i> (Crossland, 1924)		est		expN	expN		inv	exp	
<i>Ficopomatus enigmaticus</i> (Fauvel, 1923)			cas	expS	expS				est
<i>Hydroides operculata</i> (Treadwell, 1929)	est	est	est	est	inv		est/inv	est	est
<i>Leodice antennata</i> Savigny in Lamarck, 1818			exp	expS					est
<i>Leonnates indicus</i> Kinberg, 1866			exp	cas	cas				est
			exp	expS	expS				est

Species	AL	BG	GR	HR	IT	ME	RO	SL	TR
<i>Leonnates persicus</i> Wesenberg-Lund, 1949			cas		expS				est
<i>Notomastus mossambicus</i> (Thomassin, 1970)			exp		exp				est
<i>Polydora cornuta</i> Bosc, 1802	exp	est	cas	exp	est	exp	inv	exp	est
<i>Prionospio krusadensis</i> Fauvel, 1929			exp		expS				est
<i>Prionospio saccifera</i> Mackie & Hartley, 1990			unk		expS				est
<i>Pseudonereis anomala</i> Gravier, 1900	exp		est		est				est
<i>Pseudopolydora paucibranchiata</i> (Okuda, 1937)	exp	exp	est	exp	est	exp	exp	exp	est/inv
<i>Spirobranchus kraussii</i> (Baird, 1865)			exp		expS				est
<i>Streptosio gynobranchiata</i> Rice & Levin, 1998		exp			expN		inv		est
<i>Amathia verticillata</i> (delle Chiaje, 1822)	exp	exp	est/inv	est	est	exp	exp	unk	unk
<i>Diadumene lineata</i> (Verrill, 1869)		est		exp	est	exp	exp	unk	est
<i>Clytia hummelincki</i> (Leloup, 1935)	exp		exp	unk	est	exp		exp	exp
<i>Clytia linearis</i> (Thorneley, 1900)	exp		est	exp	est	exp		exp	est
<i>Garveia franciscana</i> (Torrey, 1902)	exp	est	exp	exp	est	exp	exp	exp	exp
<i>Macrorhynchia philippina</i> Kirchenpauer, 1872			exp		expS				est
<i>Cassiopea andromeda</i> (Forskål, 1775)	exp		est	expS	cas	exp			est
<i>Phyllorhiza punctata</i> Lendenfeld, 1884			est		est				est
<i>Rhopilema nomadica</i> Galil, Spanier & Ferguson, 1990			cas		cas				est
<i>Grandidierella japonica</i> Stephensen, 1938		exp		exp	est		exp	exp	exp
<i>Megabalanus tintinnabulum</i> (Linnaeus, 1758)	exp	exp	est	unk	est	exp	exp	unk	exp
* <i>Heterosaccus dollfusi</i> Boschma, 1960			exp		expS				est
* <i>Mytilicola orientalis</i> Mori, 1935		exp			expN		exp		
<i>Oithona davisae</i> Ferrari & Orsi, 1984		est			exp		est		est
<i>Pseudodiaptomus marinus</i> Sato, 1913			exp		est		exp	est	
<i>Atergatis roseus</i> (Rüppell, 1830)			est	exp	expS				est
<i>Callinectes sapidus</i> Rathbun, 1896			est		est				est
<i>Charybdis japonica</i> (A. Milne-Edwards, 1861)	est	est	est	est	est	est	est	exp	exp
<i>Charybdis longicollis</i> Leene, 1938	exp	exp	exp	exp	cas	exp	exp	exp	exp
<i>Eriocheir sinensis</i> H. Milne Edwards, 1853		exp	est	exp	expS				est
<i>Hemigrapsus sanguineus</i> (De Haan, 1835)		exp	exp	exp	cas		cas	exp	exp
* <i>Hemigrapsus takanoi</i> Asakura & Watanabe, 2005		exp		exp	expN		exp	exp	exp
<i>Ixa monodi</i> Holthuis & Gottlieb, 1956			est		expS				est
<i>Leptochela pugnax</i> de Man, 1916			exp		expS				est

Species	AL	BG	GR	HR	IT	ME	RO	SL	TR
<i>Manuta victor</i> (Fabricius, 1781)			exp		expS				est
<i>Metapenaeopsis aegyptia</i> Galil & Golani, 1990			est		expS				est
<i>Metapenaeopsis mogiensis consobrina</i> (Nobili, 1904)			est		expS				est
<i>Metapenaeus monoceros</i> (Fabricius, 1798)			exp		expS				est
<i>Metapenaeus stebbingi</i> Nobili, 1904			exp		expS				est
<i>Myra subgranulata</i> Kossmann, 1877			est		expS				est
<i>Ogyrides mjoebergi</i> (Balss, 1921)			exp		expS				est
<i>Penaeus aztecus</i> Ives, 1891			inv		est	cas			inv
<i>Penaeus japonicus</i> Spence Bate, 1888	unk	exp			cas		exp		unk
<i>Penaeus pulchricaudatus</i> Stebbing, 1914			est		expS				est
<i>Penaeus semisulcatus</i> De Haan, 1844 [in De Haan, 1833-1850]			exp		exp				est
<i>Pernon gibbesi</i> (H. Milne Edwards, 1853)	est		inv	est	est	exp		exp	est
<i>Portunus segnis</i> (Forskål, 1775)	exp		est		est				est
<i>Rhithropanopeus harrisi</i> (Gould, 1841)		est/inv			est		inv		exp
<i>Trachysalambria palaestinis</i> (Steinitz, 1932)			est		cas				est
<i>Erugosquilla massavensis</i> (Kossmann, 1880)			est		cas				est
<i>Beroe ovata</i> Bruguère, 1789		est/inv	cas		unk		inv	cas	est
<i>Mnemiopsis leidyi</i> A. Agassiz, 1865	exp	est/inv	est	est	est/inv	exp	inv	est	est
<i>Anadara kagoshimensis</i> (Tokunaga, 1906)	est	est/inv	exp	est	est/inv	exp	inv	est	est
<i>Anadara transversa</i> (Say, 1822)	exp		est	est	est/inv	exp		est	est
<i>Arcuatula senhousia</i> (Benson, 1842)	unk	exp		est	est/inv	unk	cas	inv	est
<i>Brachidontes pharaonis</i> (P. Fischer, 1870)	est		est	unk	est			cas	est
<i>Chama pacifica</i> Broderip, 1835			est		expS				est
<i>Clementia papyracea</i> (Gmelin, 1791)			cas		expS				est
<i>Dendostrea</i> cf. <i>folium</i> (Linnaeus, 1758)	unk		est		expS				est
<i>Fulvia fragilis</i> (Forskål in Niebuhr, 1775)	est		est		est				est
<i>Gafrarium savignyi</i> (Jonas, 1846)			exp		expS				est
<i>Goniobranchus annulatus</i> (Eliot, 1904)			est		expS				est
<i>Gouldiopa constemans</i> (Oliver & Zuschin, 2001)			exp		expS				cas
<i>Magallana/Crassostrea</i> sp./spp.	unk	exp	est	est	est/inv	unk	cas	inv	est
<i>Matteus regula</i> (Forskål in Niebuhr, 1775)			est		expS				est
<i>Mya arenaria</i> Linnaeus, 1758		est/inv	est	exp	est		inv	exp	est
<i>Paratapes textilis</i> (Gmelin, 1791)			exp		expS				cas

Species	AL	BG	GR	HR	IT	ME	RO	SL	TR
<i>Pinctada imbricata radiata</i> (Leach, 1814)	est		inv	est	est	est		exp	est
<i>Ruditapes philippinarum</i> (Adams & Reeve, 1850)	est	exp	exp	est	inv	cas	exp	est	est
<i>Spondylus spinosus</i> Schreibers, 1793			cas		expS				est
<i>Xenostrobus securis</i> (Lamarck, 1819)		exp			est		exp		exp
<i>Bulla arabica</i> Malaquias & Reid, 2008			est		expS				est
<i>Bursatella leachii</i> Blainville, 1817	est		est	est	est	est		est	est
<i>Conomurex persicus</i> (Swainson, 1821)	unk		est		expS	exp			est
<i>Corambe (Doridella) obscura</i> (A. E. Verrill, 1870)		est	exp		expN		inv		
<i>Diodora ruppellii</i> (G. B. Sowerby I, 1835)			exp		expS				est
<i>Finella pupoides</i> A. Adams, 1860			exp		expS				est
<i>Melibe viridis</i> (Kelaart, 1858)	exp		est	est	est	est		cas	est
* <i>Murex forskoehlII</i> Röding, 1798			exp		expS				exp
* <i>Ocenebrellus inornatus</i> (Récluz, 1851)		exp			expN		exp		exp
<i>Pseudominolia nedyma</i> (Melvill, 1897)			exp		expS				est
<i>Pyrrunculus fourierii</i> (Audouin, 1826)			cas		expS				est
<i>Rapana venosa</i> (Valenciennes, 1846)	unk	inv	cas		est		inv	est	est
<i>Rhinoclavis kochi</i> (Philippi, 1848)			cas		expS				est
<i>Sepioteuthis lessoniana</i> Férussac [in Lesson], 1831			est		expS				est
<i>Syrnola fasciata</i> Jickeli, 1882			est		expS				est
* <i>Aquilonastra burtoni</i> (Gray, 1840)			exp						exp
<i>Diadema setosum</i> (Leske, 1778)			est		expS				est
<i>Ophiactis savignyi</i> (Müller & Troschel, 1842)			cas		expS				est
<i>Ascidiaella aspersa</i> (Müller, 1776)	exp	exp	est	exp	est	exp	exp	est	est
<i>Botrylloides giganteum</i> (Pérès, 1949)			exp		unk				exp
* <i>Botrylloides nigrum</i> Herdman, 1886	exp		exp	expS	expS				exp
<i>Ciona robusta</i> Hoshino & Tokioka, 1967	exp	exp	exp	exp	est	exp	exp	exp	est
<i>Diplosoma listerianum</i> (Milne Edwards, 1841)	exp	exp	est	exp	est	exp	exp	exp	est
* <i>Ecteinascidia thurstoni</i> Herdman, 1890	exp		exp	exp	exp	exp		exp	exp
<i>Herdmania momus</i> (Savigny, 1816)			est		expS				est
<i>Microcosmus exasperatus</i> Heller, 1878			exp		expS				est
<i>Microcosmus squamiger</i> Michaelsen, 1927		exp	exp		est		exp		exp
<i>Phallusia nigra</i> Savigny, 1816			est		expS				est
* <i>Polyclinum constellatum</i> Savigny, 1816			exp		expS				exp

Species	AL	BG	GR	HR	IT	ME	RO	SL	TR
* <i>Rhodosoma turcicum</i> (Savigny, 1816)			exp		expS				exp
<i>Styela clava</i> Herdman, 1881	exp	exp	exp	exp	exp	exp	exp	exp	unk
<i>Styela plicata</i> (Lesueur, 1823)	exp	exp	exp	est	est	exp	exp	est	exp
<i>Symplegma brakenhielmi</i> (Michaelsen, 1904)			est		expS				est
<i>Apogonichthyoides pharaonis</i> (Bellotti, 1874)			est		expS				est
<i>Atherinomorus forskalii</i> (Rüppell, 1838)	exp		est		expS				est
<i>Cheilodipterus novemstriatus</i> (Rüppell, 1838)			est		expS				est
<i>Cynoglossus sinusarabici</i> (Chabanaud, 1931)			exp		expS				est
<i>Decapterus russelli</i> (Rüppell, 1830)			exp		expS				est
<i>Etrumeus golani</i> DiBattista, Randall & Bowen, 2012			est		cas				est
<i>Fistularia commersonii</i> Rüppell, 1838			inv	cas	est	cas		exp	inv
<i>Jaydia smithi</i> Kotthaus, 1970	exp		exp		expS				est
<i>Lagocephalus scleratus</i> (Gmelin, 1789)	exp		inv	est	est	cas		exp	inv
<i>Lagocephalus suezensis</i> Clark & Gohar, 1953			est		expS				est
<i>Liza carinata</i> (Valenciennes, 1836)			cas	unk	expS				est
<i>Parexocoetus mento</i> (Valenciennes, 1847)	cas		est		expS				est
<i>Penpheris rhomboidea</i> Kossmann & Räuber, 1877			est		expS				est
<i>Planiliza haematocheila</i> (Temminck & Schlegel, 1845)		est	est		expS		inv		est
<i>Plotosus lineatus</i> (Thunberg, 1787)			exp		expS				est
<i>Pteragogus trispilus</i> Randall, 2013			est		expS				est
<i>Pterois miles</i> (Bennett, 1828)	exp		est/inv		cas	exp			est/inv
<i>Sargocentron rubrum</i> (Forskål, 1775)			est		expS				est
<i>Saurida lessepsianus</i> Russell, Golani & Tikochinski, 2015	cas		est		cas				est
<i>Scomberomorus commerson</i> (Lacepède, 1800)			est		expS				est
<i>Siganus luridus</i> (Rüppell, 1829)			inv	est	est	cas		inv	inv
<i>Siganus rivulatus</i> Forskål & Niebuhr, 1775			inv	cas	cas			inv	inv
<i>Sillago suezensis</i> Golani, Fricke & Tikochinski, 2013			exp		expS				est
<i>Sphyaena chrysoaenia</i> Klunzinger, 1884			est	cas	expS				est
<i>Sphyaena flavicauda</i> Rüppell, 1838	exp		est		expS				est
<i>Stephanolepis diaspros</i> Fraser-Brunner, 1940			est	exp	est	cas		cas	est
<i>Upeneus moluccensis</i> (Bleeker, 1855)			est		cas				est
<i>Upeneus pori</i> Ben-Tuvia & Golani, 1989			est		cas				est

ing: 2), whereas *P. miles* was the only species that had solely been introduced through the Suez Canal (Table A, on-line supplement). There is an expected likelihood of all species to further expand their distribution in the ESENIAS countries, either unaided or through shipping, a possibility that in some cases is medium to high (Table 2). Out of the ten selected species, six have already impacts on the ESENIAS countries, mainly on ecosystems/biodiversity and human health (Table 2), yet they are all expected to have new/additional impacts (Table 2), as in other invasive areas, that should be addressed/mitigated to the best possible extent.

Alexandrium monilatum (J.F.Howell) Balech (Fig. 2)

Alexandrium monilatum is a thecate, chain-forming dinoflagellate. It is a coastal and estuarine planktonic species of warm temperate and tropical environments (TAYLOR *et al.*, 1995; STEIDINGER & TANGEN, 1996). The species is found in the Atlantic littoral of the USA as follows: Gulf of Mexico (CONNELL & CROSS, 1950; RAY & ALDRICH, 1967); Florida (HOWELL, 1953); Chesapeake Bay (MAY *et al.*, 2010). in the Caribbean Sea: Venezuela (HALIM, 1967), and in the tropical Pacific Ocean off Ecuador (BALECH, 1995). It was reported for the first time at the Bul-

garian coast of the Black Sea in 1991 (MONCHEVA *et al.*, 1995), where it was also later observed (MONCHEVA *et al.*, 2001; NESTEROVA *et al.*, 2008). In 2001, the species has been detected also in Odessa port, Ukraine (ANONYMOUS, 2015).

Alexandrium monilatum has been associated with bloom formation (HOWELL, 1953; PERRY *et al.*, 1979; HARDING *et al.*, 2009) and fish kills due to ichthyotoxins (GATES & WILSON, 1960; RAY & ALDRICH, 1967). Whole cells and crude extracts of *A. monilatum* have been shown experimentally to be lethal to mice, rats, fish, shellfish and cockroaches (GATES & WILSON, 1960; ALDRICH *et al.*, 1967; CLEMONS *et al.*, 1980a, b; ERKER *et al.*, 1985; MAY *et al.*, 2010). It was reported that *A. monilatum* produce PSP toxins (HSIA *et al.*, 2006), yet, to date, the species has not been related to toxic events in the Black Sea. It has been reported in blooming concentration along the Bulgarian Black Sea coast (MONCHEVA *et al.*, 1995, 2001). Blooming species cause water discoloration, especially in late summer, as well as community changes (MONCHEVA & KAMBURSKA, 2002; MONCHEVA *et al.*, 1995, 2001; VELIKOVA *et al.*, 1999). Other socio-economic effects include beach water aesthetics with a negative impact on recreation (KATSANEVAKIS *et al.*, 2014).



Fig. 2. *Alexandrium monilatum* (J.F.Howell) Balech (left; figure from WALKER & STEIDINGER (1979)) and its distribution in the Black Sea (TRICHKOVA *et al.*, 2017)

Table 2. Selected features used for horizon scanning and risk assessment, for the ten species that fact sheets were compiled within the ESENIAS-Tools project. AL=Albania, BG=Bulgaria, GR=Greece, HR=Croatia, IT=Italy, ME=Montenegro, RO=Romania, SL=Slovenia, TR=Turkey. VEC= potential vector of expansion, S=shipping, U=unaided, RoS= risk of spread, M=Medium, H=high, L=low

Species	Presence in ESENIAS countries			RoS	densities/population explosions	Impacts in ESENIAS countries	
	present	expected	VEC			current	potential
<i>Alexandrium monilatum</i>	BU	HR, IT, RO	S	M	blooms in the Black Sea	impact on ecosystem (consumption of nutrients, light decrease)	toxic event; ecosystem degradation
<i>Bonnemaisonia hamifera</i>	GR, IT, TR	AL, BG, HR, ME, RO	S, U	M/H		impact on ecosystem (growth of other algal species)	
<i>Streblospio gynobranchiata</i>	RO, TR	BG, IT	S	M	high densities in the Romanian coasts and in Izmir Bay	impact on ecosystem (alteration of indigenous biocoenoses)	invasion in polluted areas causing ecosystem degradation
<i>Cassiopea andromeda</i>	GR, IT, TR	AL, HR, ME	U	L	high densities in Greece and Turkey	impact on human health	impact on biodiversity (through predation) and ecosystem services (e.g. tourism)
<i>Oithona davisae</i>	BG, RO, TR	IT	S	L	established populations in the Black Sea	replacement of native species (e.g. <i>Oithona nana</i>)	effects on the abundance of flagellates and bacteria
<i>Penaeus aztecus</i>	GR, IT, ME, TR		U	M	dense exploitable populations in Greece and Turkey	none to date	effects on biodiversity (competition with the native <i>Penaeus kerathurus</i>)
<i>Mya arenaria</i>	BU, GR, IT, RO, TR	HR, SL	U	H	high densities in the Black Sea	invasive in the Black Sea, causing regime shifts and changes in the native communities	invasive in other areas of its distribution causing effects on the ecosystem
<i>Diadema setosum</i>	GR, TR	IT	U	M	spreading rapidly in Greece	none to date	human health; biodiversity (through predation) and ecosystem services (e.g. tourism)
<i>Microcosmus exasperatus</i>	TR	GR, IT	S, U	M	unknown	none to date	effects on biodiversity (biofouling and competition with other sedentary species)
<i>Pterois miles</i>	GR, IT, TR	AL, ME	U	M	increasing populations in Greece	none to date	human health; biodiversity (through predation)

Bonnemaisonia hamifera Hariot (Fig. 3)

Bonnemaisonia hamifera is a red alga with a heteromorphic cycle with alternation between a large, coarsely branched gametophyte and a delicate filamentous sporophyte, originally described as a separate species (*Trailliella intricata* Batters). Native in the North West Pacific, *B. hamifera* was first recorded in Europe in the British Isles (Dorset) in 1890. The species was probably introduced by shipping. Nowadays, it is widely distributed on southern and western coasts to Shetland Isles, and can be abundant in certain locations, notably where there are large, lagoon-like lower intertidal pools. In the Mediterranean Sea, the first record was as “*Trailliella intricata*” in 1909 from Tunisia. The gametophytic phase seems to have been observed only in Spain, close to Gibraltar and in Catalonia, whereas the sporophytic phase has been recorded all around the basin (VERLAQUE *et al.*, 2015). *Bonnemaisonia hamifera* can be found on rocks and on various benthic organisms.

In the north-eastern Atlantic, *B. hamifera* can have a negative impact as an ecosystem engineer by forming dense epiphytic growth on host algae (JOHANSSON *et al.*, 1998), thus reducing light and nutrient availability for those algae, in addition to preventing competing algae to colonize (SVENSSON *et al.*, 2013). By its high ratio of surface to volume, it has a greater potential for rapid uptake of nutrients in comparison to their host algae (LITTLER & LITTLER, 1980).

Native herbivores strongly preferred native algae to *B. hamifera*. *Bonnemaisonia hamifera* produces chemical grazer deterrents, mainly the secondary metabolite 1,1,3,3-tetrabromo-2-heptanone not known from the native algae of the invaded area and the importance of the chemical defence was underlined by the feeding preference of herbivores for *B. hamifera* individuals with an experimentally depleted content of 1,1,3,3-tetrabromo-2-heptanone (ENGE *et al.*, 2012). Herbivores used *B. hamifera* as a refuge for fish predation. As a result, the presence of herbivores decreases the performance of neighbouring native algae and increases growth and relative abundance of *B. hamifera* (ENGE *et al.*, 2013). The 1,1,3,3-tetrabromo-2-heptanone also works as an allelopathic compound that prevents settlement of epiphytic organisms and that can be transferred from *B. hamifera* to a native host algal species by direct contact, with an active and unaltered function, i.e. in inhibiting recruitment of native competitors (SVENSSON *et al.*, 2013). The secondary metabolite affects the natural fouling community by altering the composition, and changed the diversity by increasing the evenness and decreasing the density, indicating a broad specificity of this metabolite against bacterial colonization (e.g. PERSSON *et al.*, 2011).

Optimal experimental conditions for the biomass production of the sporophytic phase are represented by a combination of temperatures of 15-20°C, photon irradiances of 20-30 $\mu\text{mol photons m}^{-2}\text{s}^{-1}$ and long daylengths (16:8 h L:D) (NASH *et al.*, 2005).



Fig. 3. *Bonnemaisonia hamifera* Hariot (left; Photo: Marjan Richter) and its distribution in the Mediterranean Sea (TRICHKOVA *et al.*, 2017). Scale bar=1 cm



Fig. 4. *Cassiopea andromeda* (Forsskål, 1775) (left; Photo: Cemal Turan) and its distribution in the Mediterranean Sea (TRICHKOVA *et al.*, 2017; modified)

Cassiopea andromeda (Forsskål, 1775) (Fig. 4)

Cassiopea andromeda is a globally distributed in warm coastal regions semi-sessile, planktonically dispersed scyphomedusa, typically found in shallow lagoons, intertidal sand or mud flats, and around mangroves in Florida, and in the Caribbean. Its native distribution extends from the West Indian Ocean to West Pacific Ocean, and the tropical Atlantic (HOLLAND *et al.*, 2004). The presence of *C. andromeda* in the Mediterranean Sea dates prior to 1900 (MAAS, 1903).

Cassiopea andromeda is venomous (PALOMARES & PAULY, 2016) and can kill smaller fish and other marine organisms with which they come into contact, and as such is a threat to the ecosystem functioning. The species might cause an imbalance in a closed area like a bay or lagoon, but its motility is very limited, and is unlikely to spread very much.

Cassiopea andromeda is not highly dangerous because its sting is very mild. Although there has not been any record on hospitalized events of *C. andromeda* stings, this jellyfish stinging cell can possibly cause discomfort on thin or sensitive skin, as well as the eyes and lips. Moreover, *C. andromeda* do not normally swim in the water but lie on the bottom with their umbrella facing and touching the substratum and their arms, which are short and shrubby, pointing towards the surface. Swimming over the jellyfish (especially using swim fins) may cause the transparent, essentially invisible, sheets of the mucus to rise in the water

column. The stings, appearing in the form of a red rash-like skin irritation, are known for being extraordinarily itchy.

Due to its appearance and nature it may bring people snorkelling or diving into the areas in which the jelly lives, adding to the local economy.

Cassiopea andromeda feeds on copepods, cladocerans, mollusc larvae and on pelagic fish eggs and larvae (PALOMARES & PAULY, 2016). The predation upon fish larvae and eggs could potentially cause impacts on fisheries, especially in the case of consumption of commercial species, yet this is a hypothesis that needs to be further investigated.

No risk assessment analysis on biodiversity, ecosystem services or economy has been conducted. However, its presence in Turkish and Greek coastal areas that are favourite tourist destinations may lead to significant loss of ecosystem services.

Mya arenaria Linnaeus, 1758 (Fig. 5)

Mya arenaria is a bivalve that lives burrowed in muddy, sandy and gravelly sediments. It adapts well to a wide range of salinities and temperatures, and can survive both in pristine and disturbed/polluted habitats, including oxygen depleted environments, down to ≥ 200 m deep.

Archaeological evidence seems to suggest that *Mya arenaria* originated in the Pacific Ocean during the Miocene, and extended its range in the early Pliocene to the Atlantic.



Fig. 5. *Mya arenaria* Linnaeus, 1758 (left; Photo: Edoardo Turolla) and its distribution in the Mediterranean and Black Seas (TRICHKOVA *et al.*, 2017; modified)

However, the Pacific and European populations became extinct during the Pleistocene glaciations, leaving only the Northwest Atlantic population alive. *Mya arenaria* only recolonized both areas in historical times, and now occurs in a wide North East Atlantic area from the Barents Sea to Portugal, including the Baltic Sea (STRASSER, 1999; CROSS *et al.*, 2016; LASOTA *et al.*, 2016). It was also recently sighted in the Mediterranean and Black Sea, including the Turkish Straits (Bosphorus and Marmara Sea), where its presence is confirmed in five ESENIAS countries (Fig. 5). There are no convincing records from the Dardanelles and the Turkish Black Sea so far, presumably due to lack of field studies.

Mya arenaria local abundances and establishment success vary within the different ESENIAS countries. It is established in the Mediterranean Sea and the Turkish Straits, but with very low abundances and/or restricted ranges except France, where the species spread and become abundant in the Gulf of Lion (ZIBROWIUS, 2002; CROCI & TUROLLA, 2011). In the Black Sea, soon after its first record from Ukraine (1966) (BESHEVLY & KALYAGIN, 1967), the species established and became dominant in the north western and western parts of the Black Sea, and in the Sea of Azov (ZOLOTAREV, 1996). On the Bulgarian shelf, *M. arenaria* is widely distributed on sandy sediments in low salinity waters and at some sites it reaches densities of 4860 individuals m^{-2} (MARINOV, 1990). In the north western part of the Black Sea it is especially numerous in the coastal zones on muddy sediments, where its biomass exceeds 1 kg m^{-2} (ZOLOTAREV, 1996).

Mya arenaria impacts vary within the different ESENIAS countries. No impacts have been reported for the Mediterranean and the Turkish Straits, but the taxon shows invasive properties in the Black Sea dominating the soft substratum communities, causing regime shifts and structural changes in native communities/invaded habitats, and affecting sediment and water-column characteristics (KATSANEVAKIS *et al.*, 2014).

In its native area, *M. arenaria* is considered a delicacy, and is harvested by commercial fishery for culinary purposes. On the contrary, to date, the species is not commercially exploited in Europe. It is also a food source for migrating shorebirds and for a wide number of sea inhabitants (fish, sandworms and crabs, among others) (COHEN, 2005). As a suspension feeder, it plays a crucial role in filtering and cleaning water sources, and can be used as a tool in regulating and enforcing pollution standards in water quality control. It sequesters carbon in the form of calcium carbonate used for shell creation (KATSANEVAKIS *et al.*, 2014).

Streblospio gynobranchiata Rice & Levin, 1998 (Fig. 6)

Streblospio gynobranchiata is a shallow-water tube-dwelling polychaete species, found in muddy sediments of estuaries, coastal areas and harbours, at depths ranging from 0.5 to 35.6 m, where it forms dense aggregations. CINAR *et al.* (2005b) reported densities of up to 34,740 individuals m^{-2} . It prefers the upper layer of the muddy sediments with high levels of organic enrichment, rich in hydrogen sulphide and



Fig. 6. *Streblospio gynobranchiata* Rice & Levin, 1998 (left; Photo: Natalya A. Boltachova) and its distribution in the Mediterranean and Black Seas (TRICHKOVA *et al.*, 2017)

organic nitrogen (RADASHEVSKY & SELIFONOVA, 2013; BOLTACHOVA *et al.*, 2015). The tolerance of *S. gynobranchiata* to low concentrations of dissolved oxygen contributes to the colonization of substrates in severely polluted and physically degraded environments. *Streblospio gynobranchiata* is considered as one of the most successful pioneer and opportunistic species.

The native distribution area of this species is the western Atlantic (RICE & LEVIN, 1998; RADASHEVSKY & SELIFONOVA, 2013). It is one of the worst invasive species in the Mediterranean basin, where it was probably introduced via ships ballast waters (ÇINAR *et al.*, 2005a, b). It was first mentioned in the ESENIAS geographic area from the Aegean Turkish littoral (Izmir Bay), where it has been established since 2000 (ÇINAR *et al.*, 2005a, b; 2006; DAĞLI *et al.*, 2011). This species established large and dense populations in the Sea of Marmara (Istanbul area, Golden Horn Estuary, Bosphorus Strait; ÇINAR *et al.*, 2009). It was also reported in the Black Sea (MURINA *et al.*, 2008; BOLTACHOVA, 2008; RADASHEVSKY & SELIFONOVA, 2013; BOLTACHOVA *et al.*, 2015; TEACĂ *et al.*, 2015) and the Caspian Sea (TAHERI *et al.*, 2008). Recently, in 2015, this species was mentioned from the north-eastern part of the Azov Sea, from Taganrog Bay and from the Don River Delta (SYOMIN *et al.*, 2017).

To date, the impact in the ESENIAS area is a local one, with populations establishing especially in harbour areas and polluted or degraded habitats. The dense populations of the polychaete change the indigenous biocoenoses, especially in polluted soft-bottom benthic habitats (ÇINAR *et al.*, 2005b, 2011).

Dense populations and high biomass of *S. gynobranchiata* and the position it holds in the food-web as a deposit feeder in areas highly polluted with organic matter have an obvious ecological impact. In such habitats, *S. gynobranchiata* prove to be a dominant species, as well as a pollution indicator. As a result of the species biology and the large ecological tolerance of *S. gynobranchiata*, harbour areas with huge traffic and narrow gulfs with muddy bottoms are the most sensitive areas where this spionid polychaete could establish and develop large and dense populations replacing native species. The only measures to avoid the establishment of this species could be maintaining low pollution and eutrophication levels and avoiding discharge of ballast waters in the coastal areas.

Oithona davisae Ferrari & Orsi, 1984 (Fig. 7)

Oithona davisae is a pelagic cyclopoid copepod which could be the most abundant mesozooplankton species in its native habitat. It is widely spread around the Black Sea: during the first years the species developed mainly in the coastal waters but now could reach also deep waters above the depth 1000 m. In the coastal and open Black Sea this new *Oithona* taxon occupied the niche of the disappeared *Oithona nana* Giesbrecht, 1893 (SHIGANOVA *et al.*, 2012).

It is widely accepted that *O. davisae* is originally endemic to the temperate coastal waters of East Asia and its occurrence in other remote regions is due to anthropogenic introduction, mainly through ballast waters (FERRARI & ORSI, 1984; NISHIDA, 1985; HIRAKAWA, 1988). *Oithona*

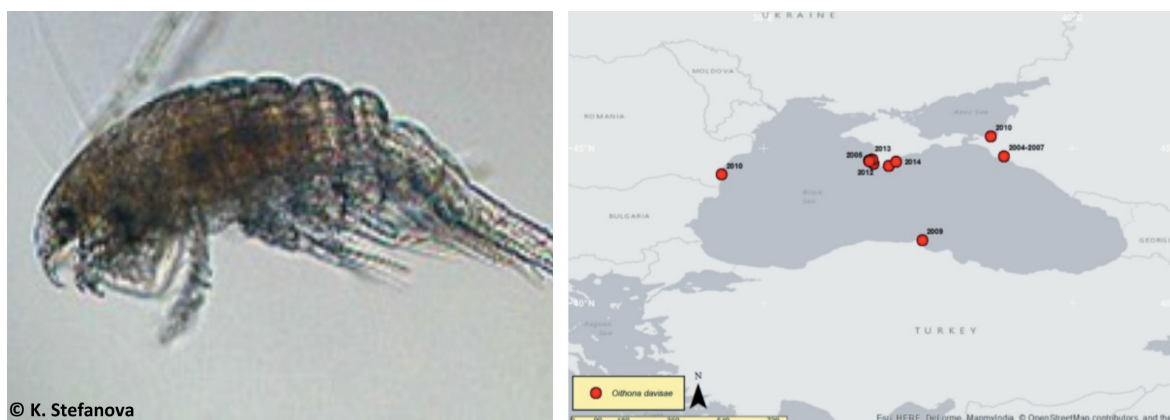


Fig. 7. *Oithona davisae* Ferrari & Orsi, 1984 (left; Photo: Kremena Stefanova) and its distribution in the Black Sea (TRICHKOVA *et al.*, 2017)

davisae usually inhabits eutrophic embayments (UYE & SANO, 1995; ALMEDA *et al.*, 2011) and is indigenous to Japan and the China Seas, and many coastal areas (RAZOULS *et al.*, 2012), but it is invasive along the USA west coast (FERRARI & ORSI, 1984) and the Spanish Mediterranean (SAIZ *et al.*, 2003). The invasive successful establishment and expansion of the thermophilic copepod *O. davisae* in the cold Black Sea is evidence of the extremely high adaptive plasticity of this species. The successful establishment of *O. davisae* to the Black Sea seems related to phytoplankton structure changes including a prevalence of small flagellates due to climate-driven effects (NESTEROVA *et al.*, 2008; MAVRODIEVA, 2012). It appears the species has expanded its distribution in the Azov Sea, as it: it has been reported in the Temruk Bay in 2010 (SVISTUNOVA, 2013).

At present, *O. davisae* successfully competes with the larger copepods *Acartia tonsa* Dana, 1849 and *Acartia clausi* Giesbrecht, 1889 in the Black Sea. It is still not clear why *O. davisae* successfully occupied the ecological niche of the disappeared *O. nana*, while the latter is not successful to return. Because of the specific features of *O. davisae*, e.g. feeding on flagellates and not on diatoms and detritus, their mass development could affect the abundance of flagellates and bacteria (ROFF *et al.*, 1995).

Penaeus aztecus Ives, 1891 (Fig. 8)

Penaeus aztecus is a shrimp species, an active swimmer and burrower (SMITHSONIAN

MARINE STATION AT FORT PIERCE, 2016), with the juveniles living also in estuarine habitats. It is found from the coastline up to 160 m deep, on muddy or sandy-muddy bottoms, sometimes mixed with sand, clay or broken shells. The adults are mostly active at night, burying in the substrate at daytime (HOLTHUIS, 1980; TAVARES, 2002).

The native distribution of the species is in the western Atlantic (PALOMARES & PAULY, 2016). *Penaeus aztecus* has also been introduced to New Caledonia and French Polynesia in the 1970s for aquaculture purposes (PALOMARES & PAULY, 2016). The first record of the species in the Mediterranean Sea was in 2009 in Antalya Bay (DEVAL *et al.*, 2010), and by 2013 the species had a well-established population in the area (GÖKOĞLU & ÖZVAROL, 2013; ÖZVAROL & GÖKOĞLU, 2014). Nowadays, *P. aztecus* has been reported from various areas throughout the Mediterranean, revealing the fast spreading of the species in the basin.

The mode of introduction of *P. aztecus* in the Mediterranean is still not clear. Some authors consider that it was most likely transferred from the USA through ballast waters (DEVAL *et al.*, 2010; KEVREKIDIS, 2014), whilst more recent discussions (CRUSCANTI *et al.*, 2015; GALIL *et al.*, 2016) suggest that it may be a result of illegal introductions (an aquaculture release/escapee). Yet, both hypotheses need further investigation.

To date, no impacts have been reported for the Mediterranean, but as the species shares the same niche and has similar biological traits with



© A. Zenetou & M. Giavasi



Fig. 8. *Penaeus aztecus* Ives, 1891 (left; Photo: Argyro Zenetos & Maria Giavasi) and its distribution in the Mediterranean Sea (TRICHKOVA *et al.*, 2017)

the indigenous commercial *Penaeus kerathurus* (Forskål, 1775) (KEVREKIDIS, 2014, 2015) the interaction amongst the two species should be monitored and assessed. In the Mediterranean, *P. aztecus* is caught with both bottom trawlers and trammel nets, along with *P. kerathurus*. Due to the low catches, it is not yet marketed nor reported separately. Nevertheless, the larger size it can attain compared to that of *P. kerathurus*, and given the fact that its presence in the catches is constantly increasing (GÖKOĞLU & ÖZVAROL, 2013), may result in its commercial exploitation in the Mediterranean Sea. Yet, the fact that *P. aztecus* is infested by the parasitic isopod *Epipe-naeon ingens* (Nobili, 1906), combined with the fact that parasitic bopyrid isopods as *E. ingens* have been found to affect its growth and reproduction (KORUN *et al.*, 2013 and references therein),

might lead to a natural control of *P. aztecus* populations. However, this hypothesis needs to be monitored and further investigated.

Diadema setosum (Leske, 1778) (Fig. 9)

Diadema setosum is a venomous sea urchin (needlespined urchin, long-spined sea urchin) with distinctively delicate hollow spines that inhabits hard substrata, rocks covered with vegetation, and coral reefs. It is widely distributed in the West Indo-Pacific, from the Red Sea, Arabian Gulf, East coast of Africa, to India, Australia and Japan at depths ranging from 0 to 70 m. In the Red Sea, Gulf of Thailand and other native regions, it is the most common sea urchin and one of the most abundant benthic invertebrates.



© A. Liami



Fig. 9. *Diadema setosum* (Leske, 1778) (left; Photo: A. Liami) and its distribution in the Mediterranean Sea (TRICHKOVA *et al.*, 2017; modified)

The tropical sea urchin fauna, which survived the Messinian crisis, but the sudden cooling that started with the Arctic glaciation 2.58 million years ago, led to the end of the tropical fauna of the Mediterranean during the third Pliocene phase (POR, 2009). Yet, *Diadema*, re-colonized the Mediterranean after an interruption of more than two million years (YOKES & GALIL, 2006). Nowadays, the species is established in the eastern Mediterranean Sea (Fig. 9).

No impact on biodiversity has been reported for the Mediterranean. It is a food source for many reef fishes. The most notable predators of adult sea urchins in the tropical Indo-Pacific are certain large reef fishes, particularly the larger wrasses, triggerfishes, puffers and porcupine fishes. In coral reef aquaria, it offers useful microhabitats to many organisms and it is considered an excellent algae controller. The risk of possible overgrazing phenomena in the ESENIAS countries deserves further study.

Diadema setosum is of medical importance. The species requires particular care when handling or working around it. Its long, slender spines may inflict painful injuries on the unwary swimmers, divers and fishermen. The spines are brittle and hollow, with barbed tips that penetrate the skin and remain imbedded in the flesh, releasing venom from their tissue and lumen. The venom may cause redness, swelling, and acute pain, which subsides after a few hours; however, spine fragments are difficult to

remove, and healing may take several weeks (YOKES & GALIL, 2006). The venom of *D. setosum* is not at all fatal to humans.

Edible and eaten in some native areas, but it is not very palatable. *Diadema setosum* is also involved in the marine aquarium trade.

Microcosmus exasperatus Heller, 1878 (Fig. 10)

Microcosmus exasperatus is a solitary ascidian with long siphons and leathery bright orange tunic with a few encrustations around the base. It typically attaches to natural and mostly artificial marine hard substrates, forming dense aggregations, which may be heavily fouled by a numerous epibionts.

Microcosmus exasperatus shows a circum-tropical distribution, being widely recorded from both the Atlantic and the Indo-Pacific regions, including the Red Sea. According to RAMOS-ESPLA *et al.* (2013) *M. exasperatus* was first recorded from Djerba Island (south of Tunisia) in 1998 and then it was observed on the coasts of Lebanon and Israel. It is commonly considered as being a Lessepsian immigrant due to multiple records from its easternmost shores. However, *Microcosmus* taxa are difficult to be identified, and therefore *M. exasperatus* worldwide identifications and possible spread patterns may be re-assessed by molecular taxonomy and phylogeography (genbank number: KT387604). Its alien Mediterranean distribution only includes



Fig. 10. *Microcosmus exasperatus* Heller, 1878 (left; Photo: Noa Shenkar) and its distribution in the Mediterranean Sea (TRICHKOVA *et al.*, 2017)

one ESENIAS country and one main sea (Turkey, Mediterranean Sea), where the taxon is only known on the basis of a single specimen (RAMOS-ESPLÁ *et al.*, 2013). Studies that are more recent have revealed that it is established in the region (ME Cinar, unpubl. Info.). However, *M. exasperatus* was recently recorded from Cyprus (GEWING *et al.*, 2016), and its known distribution may be easily concealed by taxonomic impediments and low ascidian research effort.

Native *Microcosmus* taxa are commercially exploited in the Mediterranean and consumed in France, Italy and Greece due to their strong iodine taste. The high nutritional values and the most likely similar taste of the alien taxon would suggest a possible use for culinary purposes. The typical dense populations and heavy epibiont coverage of *M. exasperatus* may contribute to its establishment as an ecosystem engineer species, as already reported for other worldwide tunicates, and may cause problems when associated with other fouling species.

Pterois miles (Bennett, 1828) (Fig. 11)

A tropical marine fish species, *Pterois miles* is a reef associated species, living depth range 25-85 m, in a variety of habitats, including natural hard bottom, artificial structures, wrecks, bridge pilings, and seagrass (FROESE & PAULY, 2016).

Outside its native range, the species is reported along the southeastern United States coast from Florida to North Carolina, from Bermuda, Bahamas, and is becoming established in

the Gulf of Mexico and the Caribbean, including northern South America, and Brazil (SCHOFIELD, 2010; FERREIRA *et al.*, 2015).

In the Mediterranean, a single record from Israel in 1991 (GOLANI & SONIN, 1992), had no follow up for about 20 years. However, a wave of *P. miles* invasion struck the eastern Mediterranean in the 2010s and is continuing at an increasing rate. In 2012 it was reported from Lebanon (BARICHE *et al.*, 2013), and a year later from Cyprus (EVRIPIDOU, 2013) where, in just one year, it had colonized nearly all of Cyprus' southeastern coast (JIMENEZ *et al.*, 2016). In the following years, the species was sighted in several Mediterranean locations (Fig. 11).

Pterois miles is native to the Indo-Pacific realm (FROESE & PAULY, 2016), and it is thought to have been introduced to the western Atlantic in the mid 1990's by aquarists (HARE & WHITFIELD, 2003). In the Mediterranean, the most likely introduction vector is considered to be the Suez Canal either through ballast water released from vessels crossing the canal (even though globally successful introductions of Scorpaenidae through ballast water have not been reported; see HARE & WHITFIELD, 2003) or by adult migration. However, secondary introductions through additional aquarium release events cannot be ruled out, and genetic connectivity studies are required to trace the origin of the Mediterranean populations. A recent genetic study presents evidence, albeit from a limited sample size, that supports the immigration of a few individuals from the Red Sea via the Suez Canal as the most



Fig. 11. *Pterois miles* (Bennett, 1828) (left; Photo: Gerasimos Kondylatos) and its distribution in the Mediterranean Sea (TRICHKOVA *et al.*, 2017; modified)

likely introduction mechanism (BARICHE *et al.*, 2017).

In the west Atlantic invaded range, lionfish have reached densities that are far higher than those reported from their native range (KULBICKI *et al.*, 2012), exhibit extraordinary predation rates and are having dramatic impacts on the Caribbean ecosystem by displacing native species and disrupting food webs (LESSER & SLATTERY 2011; ALBINS 2013).

In the Mediterranean, owing to the young age of the invasion, strong impacts on ecosystems have not been reported yet, as is the case in the invaded Atlantic distribution (for details see ALBINS & HIXON, 2008; LESSER & SLATTERY, 2011; GREEN *et al.*, 2012, 2014; ALBINS, 2013; FALETTI *et al.*, 2013; BENKWITT, 2015; BALLEW *et al.*, 2016; PALMER *et al.*, 2016) but are the subject of ongoing research. In addition, the full extent of the socio-economic damage potentially caused by lionfish is yet to be realised, since the full spectrum and intensity of its ecological impacts and interactions with native species is not fully known.

Pterois miles has a direct impact on human health. The sting from its venomous spines can cause irritation, inflammation, pain and even serious complications in the case of an allergic reaction, putting fishermen, divers and other potential stakeholders at risk. On the other hand, it appears to be very attractive to divers, many of who specifically request dives in areas where the lionfish is known to be present (JIMENEZ *et al.*, 2017), at least in the Eastern Mediterranean, where the species is still a novel sighting, similar to what was observed in the Atlantic in the early stages of the invasion. Regardless of its venomous spines, the species is a popular aquarium fish and may create an extra source of income for fishermen who capture and supply the aquarium trade. Moreover, it is consumed in subsistence fisheries in its native area and can provide an alternative fish stock and food source

in the invaded range. Its consumption is already promoted in some countries as a means of introducing control measures (NUÑEZ *et al.*, 2012).

CONCLUSIONS

This is the first effort to compile a list of marine invasive/potential invasive species in the ESENIAS countries. Such lists are an important initial step for the development of an early warning system, through a horizon scanning process (ROY *et al.*, 2015). Hence, based on the information presented here identification of future invasions in the ESENIAS countries is possible. It is essential, as a way forward for the countries referred herewith in, to further plan/enforce effective early warning and rapid response mechanisms, as well as take management measures for species that are already widely spread, in compliance with the EU Regulations (EU, 2014). Yet, such a list should be regularly revised and updated so that the management measures enforced would be effective.

ACKNOWLEDGMENTS

This study was conducted in the frame of the project entitled “East and South European Network for Invasive Alien Species - a tool to support the management of alien species in Bulgaria (ESENIAS-TOOLS)”, an EEA funded project (Contract No. Д-33-51/30.06.2015). The authors would also like to thank the following colleagues for providing photos: M. RICHTER (photo of *Bonnemaisonia hamifera*), C. TURAN (photo of *Cassiopea andromeda*), E. TUROLLA (photo of *Mya arenaria*), N.A. BOLTACHOVA (photo of *Streblospio gynobranchiata*), A. LIAMI (photo of *Diadema setosum*) and G. KONDYLATOS (photo of *Pterois miles*). Finally, special thanks to L. FILIPOV for creating the ESENIAS maps, and N.-J. XENTIDIS for the modified maps.

REFERENCES

- ALBINS, M.A. 2013. Effects of invasive Pacific red lionfish *Pterois volitans* versus a native predator on Bahamian coral-reef fish communities. *Biol. Invasions*, 15: 29-43.
- ALBINS, M.A. & M.A. HIXON. 2008. Invasive Indo-Pacific lionfish *Pterois volitans* reduce recruitment of Atlantic coral-reeffishes. *Mar. Ecol. Prog. Ser.*, 367: 233-238.
- ALDRICH, D.V., S.M. RAY & W.B. WILSON. 1967. *Gonyaulax monilata*: population growth and development of toxicity in cultures. *J. Protozool.*, 14: 636-639.
- ALMEDA, R., M. ALCARAZ, A. CALBET & E. SAIZ. 2011. Metabolic rates and carbon budget of early developmental stages of the marine cyclopoid copepod *Oithona davisae*. *Limnol. Oceanogr.*, 56(1): 403-414.
- ANONYMOUS. 2015. *Alexandrium monilatum*. In: AquaNIS. Editorial Board, 2015. Information system on Aquatic Non-Indigenous and Cryptogenic Species. World Wide Web electronic publication. <http://www.corpi.ku.lt/databases/index.php/aquanis/introductions/view/id/4053>. Accessed on 2017-06-26
- BALECH, E. 1995. The genus *Alexandrium* Halim (Dinoflagellata). Publ. Sherkin Island marine station Sherkin Island Co. Cork Ireland, 151 pp.
- BALLEW, N.G., N.M. BACHELER, G.T. KELLISON & A.M. SCHUELLER. 2016. Invasive lionfish reduce native fish abundance on a regional scale. *Sci. Rep.*, 6: 32169.
- BARICHE, M., M. TORRES & E. AZZURRO. 2013. The presence of the invasive Lionfish *Pterois miles* in the Mediterranean Sea. *Medit. Mar. Sci.*, 14(2): 292-294.
- BARICHE, M., P. KLEITOU, S. KALOGIROU & G. BERNARDI. 2017. Genetics reveal the identity and origin of the lionfish invasion in the Mediterranean Sea. *Sci. Rep.* 7: 6782. doi:10.1038/s41598-017-07326-1.
- BENKWITT, C.E. 2015. Non-linear effects of invasive lionfish density on native coral-reef fish communities. *Biol. Invasions*, 17: 1383-1395.
- BESHEVLY, L.E. & V.A. KALYAGIN. 1967. The finding of the mollusc *Mya arenaria* L. (Bivalvia) in the northwestern part of the Black Sea. *Vestnik Zool.*, 3: 82-84 (in Russian).
- BOLTACHOVA, N.A. 2008. Finding of new alien species *Streblospio gynobranchiata* Rice et Levin, 1998 (Polychaeta: Spionidae) in the Black Sea. *Mar. Ecol. J.*, 7(4): p. 12 (in Russian).
- BOLTACHOVA, N.A., E.V. LISITSKAYA & D.V. PODZOROVA. 2015. The population dynamics and reproduction of *Streblospio gynobranchiata* (Annelida, Spionidae), an alien polychaete worm, in the Sevastopol Bay (the Black Sea). *Ecol. Mont.*, 4: 22-28.
- CBD. 2014. Pathways of Introduction of Invasive Species, their Prioritization and Management. UNEP/CBD/SBSTTA/18/9/Add.1. Secretariat of the Convention on Biological Diversity, Montréal. Pages??
- ÇINAR, M.E., H. BALKIS, S. ALBAYRAK, E. DAĞLI & S.A. KARHAN. 2009. Distribution of polychaete species (Annelida: Polychaeta) on the polluted soft substrate of the Golden Horn Estuary (Sea of Marmara), with special emphasis on alien species. *Cah. Biol. Mar.*, 50: 11-17.
- ÇINAR, M.E., M. BILECENOĞLU, B. ÖZTÜRK, T. KATAGAN, & V. AYSEL. 2005a. Alien species on the coast of Turkey. *Medit. Mar. Sci.*, 6(2): 119-146.
- ÇINAR, M.E., M. BILECENOĞLU, B. ÖZTÜRK, T. KATAGAN, M.B. YOKES, V. AYSEL, E. DAĞLI, S. ACIK, T. ÖZCAN & H. ERDOĞAN. 2011. An updated review of alien species on the coasts of Turkey. *Medit. Mar. Sci.*, 12(2): 257-315.
- ÇINAR, M.E., Z. ERGEN, E. DAGLI & M.E. PETERS-EN. 2005b. Alien species of spionid polychaetes (*Streblospio gynobranchiata* and *Polydora cornuta*) in Izmir Bay, eastern Mediterranean. *J. Mar. Biol. Assoc. U.K.*, 85(4): 821-827.
- ÇINAR, M.E., T. KATAGAN, B. ÖZTÜRK, Ö. EGEMEN, Z. ERGEN, A. KOCATAS, M. ÖNEN, F. KIRKIM, K. BAKIR, G.KURT, E. DAĞLI, A. KAYMAKÇI, S. AÇIK, A. DOĞAN & T. ÖZCAN. 2006. Temporal changes of soft-bottom zoobenthic communities in and around Alsancak Harbor (Izmir

- Bay, Aegean Sea), with special attention to the autecology of exotic species, *Mar. Ecol.*, 27: 229-246.
- CLEMONS, G.P., D.V. PHAM & J.P. PINION. 1980a. Insecticidal activity of *Gonyaulax* (Dinophyceae) cell powders and saxitoxin to the German cockroach. *J. Phycol.*, 16: 305-307
- CLEMONS, G.P., J.P. PINION, E. BASS, D.V. PHAM, M. SHARIF & J.G. WUTOH. 1980b. A hemolytic principle associated with the red tide dinoflagellate *Gonyaulax monilata*. *Toxicon*, 18: 323-326.
- COHEN, A.N. 2005. Guide to the exotic species of San Francisco Bay. San Francisco Estuary Institute, Oakland, California. Available at: www.exoticguide.org LAST ACCESSED - DATE
- CONNELL, C.H. & J.B. CROSS. 1950. Mass mortality of fish associated with the protozoan *Gonyaulax* in the Gulf of Mexico. *Science*, 112: 359.
- CROCETTA, F. & E. TUROLLA. 2011. *Mya arenaria* Linné, 1758 (Mollusca: Bivalvia) in the Mediterranean: its distribution revisited. *J. Biol. Res.-Thessaloniki*, 16: 188-193
- CROSS, M.E., C.R. BRADLEY, T.F. CROSS, S. CULLOTY, S. LYNCH, P. MCGINNITY, R.M. O'RIORDAN, S. VARTIA & P.A. PRODÖHL. 2016. Genetic evidence supports recolonisation by *Mya arenaria* (L.) of Western Europe from North America. *Mar. Ecol. Prog. Ser.*, 549: 99-112.
- CRUSCANTI, M., G. INNOCENTI, J.R. ALVARADO BREMER & B.S. GALIL. 2015. First report of the brown shrimp *Penaeus aztecus* Ives, 1891 (Crustacea, Decapoda, Penaeidae) in the Tyrrhenian Sea. *Mar. Biodiv. Records*, 8: 1-4.
- DAĞLI, E., M.E. ÇINAR & Z. ERGEN. 2011. Spionidae (Annelida: Polychaeta) from the Aegean Sea (eastern Mediterranean). *Ital. J. Zool.*, 78(S1): 49-64.
- DEVAL, M.C., Y. KAYA, O. GÜVEN, M. GÖKOĞLU & C. FROGLIA. 2010. An unexpected find of the western Atlantic shrimp, *Farfantepenaeus aztecus* (Ives, 1891) (Decapoda, Penaeidae) in Antalya bay, Eastern Mediterranean Sea. *Crustaceana*, 83(12): 1531-1537.
- ENGE, S., G.M. NYLUND, T. HARDER & H. PAVIA. 2012. An exotic chemical weapon explains low herbivore damage in an invasive alga. *Ecology*, 93(12): 2736-2745.
- ENGE, S., G.M. NYLUND & H. PAVIA. 2013. Native generalist herbivores promote invasion of a chemically defended seaweed via refuge mediated apparent competition. *Ecol. Lett.*, 16(4): 487-492.
- ERKER, E.F., L.J. SLAUGHTER, E.L. BASS, J. PINION & J. WUTOH. 1985. Acute toxic effects in mice of an extract from the marine algae *Gonyaulax monilata*. *Toxicon*, 23: 761-767.
- ESSL, F., S. BACHER, T.M. BLACKBURN, O. BOOY, G. BRUNDU, S. BRUNEL, A.-C. CARDOSO, R. ESCHEN, B. GALLARDO, B. GALIL, E. GARCÍA-BERTHO, P. GENOVESI, Q. GROOM, C. HARROWER, P.E. HULME, S. KATSANEVAKIS, M. KENIS, I. KÜHN, S. KUMSCHICK, A.F. MARTINOU, W. NENTWIG, C. O'FLYNN, S. PAGAD, J. PERGL, P. PYŠEK, W. RABITSCH, D.M. RICHARDSON, A. ROQUES, H.E. ROY, R. SCALERA, S. SCHINDLER, H. SEEBENS, S. VANDERHOEVEN, M. VILÀ, JOHN R. U. WILSON, A. ZENETOS & J.M. JESCHKE. 2015. Crossing frontiers in tackling pathways of biological invasions. *BioScience*, 65 (8): 769-782. DOI 10.1093/biosci/biv082
- EU. 2008. Directive of the European Parliament and the Council Establishing a Framework for Community Action in the Field of Marine Environmental Policy (Marine Strategy Framework Directive). European Commission. Directive 2008/56/EC, OJ L 164
- EU. 2011. Our life insurance, our natural capital: an EU biodiversity strategy to 2020. COM/2011/244, European Commission, Brussels, 16 pp.
- EU. 2014. Regulation (EU) No 1143/2014 of the European Parliament and of the 464 Council of 22 October 2014 on the prevention and management of the introduction and 465 spread of invasive alien species. Official Journal of the European Union, 57, 35.
- EVRIPIDOU, S. 2013. Toxic lionfish makes its way to Cyprus waters. <http://www.cyprus-mail.com/cyprus/toxic-Lionfish-makes-its-way-cyprus-waters/20130220>.
- FALETTI, M.E., R.D. ELLIS & F. BAY. 2013. Novel Predator, Novel Habitat : a diet analysis and

- experimental test of the ecological effects of invasive lionfish in Florida Bay. In: 66th Annual Conference, Gulf and Caribbean Fisheries Institute, 4-8 November. pp. 217-221.
- FERRARI, F. & J. ORSI. 1984. *Oithona davisae*, new species, and *Limnoithona sinensis* (Burkckhard, 1912) (Copepoda: Oithonidae) from the Sacramento - San Joaquin Estuary, California. *J. Crustac. Biol.*, 4(1): 106-126.
- FERREIRA, C.E.L., O.J. LUIZ, S.R. FLOETER, M.B. LUCENA, M.C. BARBOSA, C.R. ROCHA & L.A. ROCHA. 2015. First record of invasive lionfish (*Pterois volitans*) for the Brazilian coast. *PLoS One* 10: 1–5. doi:10.1371/journal.pone.0123002.
- FROESE, R. & D. PAULY. 2016. FishBase. World Wide Web electronic publication. www.fishbase.org, version (10/2016).
- GALIL, B.S. 2000. A Sea Under Siege – Alien Species in the Mediterranean. *Biol. Inv.*, 2(2): 177–186.
- GALIL, B.S., G. INNOCENTI, J. DOUEK, G. PAZ & B. RINKEVICH. 2016. Foul play? On the rapid spread of the brown shrimp *Penaeus aztecus* Ives, 1891 (Crustacea, Decapoda, Penaeidae) in the Mediterranean, with new records from the Gulf of Lion and the southern Levant. *Mar. Biodiv.*, 1-7. DOI 10.1007/s12526-016-0518-x
- GATES, J.A. & W.B. WILSON. 1960. The toxicity of *Gonyaulax monilata* Howell to *Mugil cephalus*. *Limnol. Oceanogr.*, 5: 171-174.
- GEWING, M.T., O. BRONSTEIN, L.R. NAGAR, I. GRANOT, O. FRID, & N. SHENKAR. 2016. First record of the non-indigenous ascidian *Microcosmus exasperatus*, Heller 1878, in Cyprus. *Mar. Biodiv.*, 46(4): 937-941.
- GÖKOĞLU, M. & Y. ÖZVAROL. 2013. Biogeographic expansion of *Farfantepenaeus aztecus* (Ives, 1891) (Decapoda: Penaeidae) in the Eastern Mediterranean Sea. p. 475. In: New Mediterranean Marine biodiversity records (December 2013). BILECENOGLU *et al.*, *Medit. Mar. Sci.*, 14(2): 463-480.
- GOLANI, D. & O. SONIN. 1992. New records of the Red Sea fishes, *Pterois miles* (Scorpaenidae) and *Pteragogus pelycus* (Labridae) from the eastern Mediterranean Sea. *Jap. J. Ichthyol.*, 39: 167-169.
- GREEN, S.J., J.L. AKINS, A. MALJKOVIC & I.M. COTE. 2012. Invasive lionfish drive Atlantic coral reef fish declines. *PLoS One*, 7(3): e32596.
- GREEN, S. J., N.K. DULVY, A.M.L. BROOKS, J.L. AKINS, A.B. COOPER, S. MILLER & I.M. COTÉ. 2014. Linking removal targets to the ecological effects of invaders: A predictive model and field test. *Ecol. Appl.* 24, 1311–1322. doi:10.1890/13-0979.1
- HALIM, Y. 1967. Dinoflagellates of the south-east Caribbean sea. *Int. Rev. gesamten Hydrobiol.*, 52(5): 701-755.
- HARDING, J.M., R. MANN, P. MOELLER & M. HSIA. 2009. Mortality of the veined rapa whelk, *Rapana venosa*, in relation to a bloom of *Alexandrium monilatum* in the York River, United States. *J. Shellfish Res.*, 28: 363–367.
- HARE, J.A. & P.E. WHITFIELD. 2003. An integrated assessment of the introduction of lionfish (*Pterois volitans* / *miles* complex) to the Western Atlantic Ocean. NOAA Tech. Memo., 21-22 pp.
- HIRAKAWA, K. 1988. New records of the North Pacific coastal planktonic copepods, *Acartia omorii* (Acartiidae) and *Oithona davisae* (Oithonidae) from southern Chile. *Bull. Mar. Sci.*, 42: 337-339.
- HOLLAND, B.S., M.N. DAWSON, G.L. CROW & D.K. HOFMANN. 2004. Global phylogeography of *Cassiopea* (Scyphozoa: Rhizostomeae): molecular evidence for cryptic species and multiple invasions of the Hawaiian Islands. *Mar. Biol.*, 145: 1119-1128.
- HOLTHUIS, L.B. 1980. FAO Species Catalogue. Shrimps and prawns of the world. An annotated catalogue of species of interest to fisheries. FAO Fisheries Synopsis, 1(125): 1-271.
- HOWELL, J.F. 1953. *Gonyaulax monilata*, sp. nov., the causative dinoflagellate of a red tide on the east coast of Florida in August–September, 1951. *Trans. Am. Microsc. Soc.*, 72: 153–156.
- HSIA, M.H., S.L. MORTON, L.L. SMITH, K.R. BEAUCHESNE, K.M. HUNCIK & P.D.R. MOE-

- LLER. 2006. Production of goniodomin A by the planktonic, chain-forming dinoflagellate *Alexandrium monilatum* (Howell) Balech isolated from the Gulf coast of the United States. *Harmful Algae*, 5: 290-299.
- JIMENEZ, C., V. ANDREOU, L. HADJIOANNOU, R.A. ALHAJJA & P. PATSALOU. 2017. Not everyone's cup of tea: Public perception of culling invasive lionfish in Cyprus. *J. Black Sea/Mediterranean Environ.*, 23: 38-47.
- JIMENEZ, C., A. PETROU, V. ANDREOU, L. HADJIOANNOU, W. WOLF, N. KOUTSOLOUKAS & R.A. ALHAJJA. 2016. Veni, Vidi, Vici: The successful establishment of the lionfish *Pterois miles* in Cyprus (Levantine Sea). *Rapp. Comm. int. Mer Médit.*, 41: p. 417.
- JOHANSSON, G., B.K. ERIKSSON, M. PEDERSEN & P. SNOEIJIS. 1998. Long-term changes of macroalgal vegetation in the Skagerrak area. *Hydrobiologia*, 385: 121-138.
- KARACHLE, P.K., A. ZENETOS & N.J. XENTIDIS. 2017. The ESENIAS countries' marine Alien Species experts: an updated inventory. *Acta Zool. Bulgarica*, suppl 9: 261-282.
- KATSANEVAKIS, S., A. ZENETOS, V. MAČIĆ, S. BEQIRAJ, D. POURSANIDIS & L. KASHTA. 2011. Invading the Adriatic: spatial patterns of marine alien species across the Ionian-Adriatic boundary. *Aquat. Biol.*, 13: 107-118.
- KATSANEVAKIS, S., I. WALLENTINUS, A. ZENETOS, E. LEPPÄKOSKI, M.E. ÇINAR, B. ÖZTÜRK, M. GRABOWSKI, D. GOLANI & A.C. CARDOSO. 2014. Impacts of invasive alien marine species on ecosystem services and biodiversity: a pan-European review. *Aquat. Invasion.*, 9: 391-423.
- KEVREKIDIS, K. 2014. The occurrence of the Atlantic penaeid prawn *Farfantepenaeus aztecus* (Ives, 1891) in the Thermaikos Gulf (Aegean Sea, Eastern Mediterranean): considerations on the potential establishment and impact on the autochthonous *Melicerus kerathurus* (Forskål, 1775). *Crustaceana*, 87: 1606-1619.
- KEVREKIDIS, K. 2015. The Atlantic shrimp *Farfantepenaeus aztecus* (Ives, 1891): an alien and commercial species in the Greek Seas (in Greek). *Hellenic Fishing News* (<http://alieftikanea.gr/i-atlantiki-garida-farfante-penaeus-aztecus-ives-1891-ena-allochthono-ke-emporiko-idos-stis-ellinikes-thalasses/>).
- KORUN, J., M. GÖKOĞLU, B.A. BALCI & Y. ÖZVAROL. 2013. Infestation of brown shrimp, *Farfantepenaeus aztecus*, Ives (1891) (Penaeidae) by *Epipenaeon ingens*, Nobili (1906) (Isopoda, Bopyridae) from the Antalya Bay, Turkey. *Rev. Médec. Vét.*, 164(12): 559-563.
- KULBICKI, M., J. BEETS, P. CHABANET, K. CURE, E. DARLING, S.R. FLOETER, R. GALZIN, A. GREEN, M. HARMELIN-VIVIEN, M. HIXON, Y. LETOURNEUR, T.L. DE LOMA, T. MCCLANAHAN, J. MCILWAIN, G. MOUTHAM, R. MYERS, J.K. O'LEARY, S. PLANES, L. VIGLIOLA & L. WANTIEZ. 2012. Distributions of Indo-Pacific lionfishes *Pterois* spp. in their native ranges: Implications for the Atlantic invasion. *Mar. Ecol. Prog. Ser.*, 446: 189-205.
- LASOTA, R., K. PIERSCIENIAK, P. GARCIA, B. SIMONBOUHET & M. WOŁOWICZ. 2016. Large-scale mitochondrial COI gene sequence variability reflects the complex colonization history of the invasive soft-shell clam, *Mya arenaria* (L.) (Bivalvia). *Estuar. Coast. Shelf Sci.*, 181: 256-265.
- LEPPAKOSKI, E. & P.E. MIHNEA. 1996. Enclosed seas under man-induced change: a comparison between the Baltic and Black Seas. *Ambio*, 25(6): 380-389.
- LESSER, M.P. & M. SLATTERY. 2011. Phase shift to algal dominated communities at mesophotic depths associated with lionfish (*Pterois volitans*) invasion on a Bahamian coral reef. *Biol. Invasions*, 13: 1855-1868.
- LITTLER, M.M. & D.S. LITTLER. 1980. The evolution of thallus form and survival strategies in benthic marine macroalgae: field and laboratory tests of a functional form model. *Am. Nat.*, 116: 25-44.
- MAAS, O. 1903. Die Scyphomedusen der Siboga-expedition [Siboga-expeditie 11.]. E.J. Brill, Leiden, viii + 91 pp. + plates I-XII
- MARCHINI A., J. FERRARIO & A. OCCHIPINTI-AMBROGI. 2013. Recent additions to the alien marine biota along Italian coasts. *Rapp. Comm. int. Mer Médit.*, 40: p. 887.
- MARINOV, T. 1990. The zoobenthos from the Bulgarian sector of the Black Sea (in Bulgarian). *Bulg. Acad. Sci. Publ.*, Sofia, 195 pp.

- MAVRODIEVA, R. 2012. Mechanisms of interactions between phytoplankton and marine environmental factors (biotic and abiotic) in the Bulgarian Black Sea coastal waters (Sozopol Bay). PhD Thesis, Institute of Biodiversity and Ecosystem Research, Bulgarian Academy of Science, Sofia, 193 pp.
- MAY, S.P., J.M. BURKHOLDER, S.E. SHUMWAY, H. HEGARET, G.H. WIKFORS & D. FRANK. 2010. Effects of the toxic dinoflagellate *Alexandrium monilatum* on survival, grazing and behavioral response of three ecologically important bivalve mollusks. *Harmful Algae*, 9: 281-291.
- MCGEOCH, M.A., P. GENOVESI, P.J. BELLINGHAM, M.J. COSTELLO, C. MCGRANNACHAN & A. SHEPPARD. 2016. Prioritizing species, pathways, and sites to achieve conservation targets for biological invasion. *Biol. Invasions*, 18: 299-314.
- MONCHEVA, S. & L. KAMBURSKA. 2002. Black Sea plankton stowaways - why a crucial issue to safeguard Black Sea biodiversity. CIESM Workshop Monographs, Monaco, no. 20, 136 pp.
- MONCHEVA, S., O. GOTSIS-SKRETAS, K. PAGOU & A. KRASTEV. 2001. Phytoplankton blooms in Black Sea and Mediterranean coastal ecosystems subjected to anthropogenic eutrophication: similarities and differences. *Estuar. Coast. Shelf S.*, 53(3): 281-295.
- MONCHEVA, S., V. PETROVA-KARADJOVA & A. PALASOV. 1995. Harmful algae blooms along the Bulgarian Black Sea coast and possible patterns of fish and zoobenthic mortalities. In: P. Lassus, G. Arzul, E. Denn and P. Gentien (Editors). *Harmful Marine Algal Blooms*. Lavoisier Publ. Inc., pp. 193-198.
- MURINA, V.V., Z.P. SELIFONOVA & V.F. MELNIK. 2008. Finding of polychaete *Streblospio* sp. (Polychaeta: Spionidae) in the Novorossiysk port of the Black Sea (in Russian). *Mar. Ecol. J.*, 7(1): p. 46.
- NASH, R., F. RINDI & M.D. GUIRY. 2005. Optimum conditions for cultivation of the *Trailliella* phase of *Bonnemaisonia hamifera* Hariot (Bonnemaisoniales, Rhodophyta), a candidate species for secondary metabolite production. *Bot. Mar.*, 48(4): 257-265.
- NESTEROVA, D., S. MONCHEVA, A. MIKAELIAN, A. VERSHININ, V. AKATOV, L. BOICENCO, Y. AKTAN, F. SAHIN & T. GVARISHVILI. 2008. The State of Phytoplankton. In: T. Oguz (Editor). *BSC, 2008. State of the Environment of the Black Sea (2001 - 2006/7)*. Publications of the Commission on the Protection of the Black Sea against pollution (BSC) 2008-3, Istanbul, pp. 173-200.
- NISHIDA, S. 1985. Taxonomy and distribution of the family Oithonidae (Copepoda, Cyclopoida) in the Pacific and Indian Oceans. *Bull. Ocean Res. Inst., University of Tokyo*, 20: 1-167.
- NUÑEZ, M. A., S. KUEBBING, R.D. DIMARCO & D. SIMBERLOFF. 2012. Invasive Species: To eat or not to eat, that is the question. *Conserv. Lett.* 5, 334-341. doi:10.1111/j.1755-263X.2012.00250.x.
- ÖZVAROL, Y. & M. GÖKOĞLU. 2014. Biological data on northern brown shrimp *Farfantepenaeus aztecus* (Ives, 1891) (Decapoda: Penaeidae) in the Eastern Mediterranean Sea. p. 214-215. In: *New Fisheries-related data from the Mediterranean Sea (April 2014)*. Stergiou *et al.*, *Medit. Mar. Sci.*, 15 (1): 213-224.
- PALMER, G., J.D. HOGAN, B.D. STERBA-BOATWRIGHT & R.D. OVERATH. 2016. Invasive lionfish *Pterois volitans* reduce the density but not the genetic diversity of a native reef fish. *Mar. Ecol. Prog. Ser.*, 558: 223-234.
- PALOMARES, M.L.D. & D. PAULY (Editors) 2016. *SeaLifeBase*. World Wide Web electronic publication. www.sealifebase.org, version (01/2016). (Accessed 04/06/2016).
- PERRY, H.M., K.C. STUCK & H.D. HOWSE. 1979. 1st record of a bloom of *Gonyaulax monilata* in coastal waters of Mississippi. *Gulf Res. Rep.*, 6: 313-316.
- PERSSON, F., R. SVENSSON, G.M. NYLUND, N.J. FREDRIKSSON, H. PAVIA & M. HERMANSSON. 2011. Ecological role of a seaweed secondary metabolite for a colonizing bacterial community. *Biofouling*, 27(6): 579-588.
- POR, F.D. 2009. Tethys returns to the Mediterranean: success and limits of tropical recolonization. *BioRisk*, 3: 5-19.
- RADASHEVSKY, V.I. & Z.P. SELIFONOVA. 2013.

- Records of *Polydora cornuta* and *Streblospio gynobranchiata* (Annelida, Spionidae) from the Black Sea. *Medit. Mar. Sci.*, 14(2): 261-269.
- RAMOS-ESPLÁ, A., A. IZQUIERDO-MUÑOZ & M.E. ÇINAR. 2013. *Microcosmus exasperatus* (Asciidiacea: Pyuridae), current distribution in the Mediterranean Sea. *Mar. Biodiv. Records*, 6, e89.
- RAY, S.M. & D.V. ALDRICH. 1967. Ecological interactions of toxic dinoflagellates and molluscs in the Gulf of Mexico. In: F.E. Russell & P.R. Saunders (Editors). *Animal Toxins, First International Symposium on Animal Toxins*, Pergamon Press, New York, pp. 75-83.
- RAZOULS, C., F. DE BOVEE, J. KOENIG & N. DESREUMAUX. 2012. Diversity and Geographic Distribution of Marine Planktonic Copepods. Available from: <http://copepodes.obs-banyuls.fr/en/fichesp.php?sp=1776>
- RICE, S.A. & L. LEVIN. 1998. *Streblospio gynobranchiata*, a new spionid polychaete species (Annelida: Polychaeta) from Florida and the Gulf of Mexico with an analysis of phylogenetic relationships within the genus *Streblospio*. *Proc. Biol. Soc. Wash.*, 111(3): 694-707.
- ROFF, J., T. TURNER, K. WEBBER & R. HOPCROFT. 1995. Bacterivory by tropical copepod nauplii: extent and possible significance. *Aquat. Microb. Ecol.*, 9(2): 165-175.
- ROY, H.E., T. ADRIAENS, D.C. ALDRIDGE, S. BACHER, J.D.D. BISHOP, T.M. BLACKBURN, E. BRANQUART, J. BRODIE, C. CARONERAS, E.J. COOK, G.H. COPP, H.J. DEAN, J. EILENBERG, F. ESSL, B. GALLARDO, M. GARCIA, E. GARCÍA-BERTHOUD, P. GENOVESI, P.E. HULME, M. KENIS, F. KERCKHOF, M. KETTUNEN, D. MINCHIN, W. NENTWIG, A. NIETO, J. PERGL, O. PESCOFF, J. PEYTON, C. PREDAL, W. RABITSCH, A. ROQUES, S. RORKE, R. SCALERA, S. SCHINDLER, K. SCHÖNRÖGGE, J. SEWELL, W. SOLARZ, A. STEWART, E. TRICARICO, S. VANDERHOEVEN, G. VAN DER VELDE, M. VILÀ, C.A. WOOD & A. ZENETOS. 2015. Invasive Alien Species - Prioritising prevention efforts through horizon scanning ENV.B.2/ETU/2014/0016. European Commission, 227 pp.
- ROY, H.E., H. HESKETH, B.V. PURSE, J. EILENBERG, A. SANTINI, R. SCALERA, G.D. STENTIFORD, T. ADRIAENS, K. BACELA-SPYCHALSKA, D. BASS, K.M. BECKMANN, P. BESSELL, J. BOJKO, O. BOOY, A.C. CARDOSO, F. ESSL, Q. GROOM, C. HARROWER, R. KLEESPIES, A.F. MARTINO, M.M. VAN OERS, E.J. PEELER, J. PERGL, W. RABITSCH, A. ROQUES, F. SCHAFFNER, S. SCHINDLER, B.R. SCHMIDT, K. SCHÖNRÖGGE, J. SMITH, W. SOLARZ, A. STEWART, A. STROO, E. TRICARICO, K.M.A. TURVEY, A. VANNINI, M. VILÀ, S. WOODWARD, A.A. WYNNS, & A.M. DUNN. 2017. Alien Pathogens on the Horizon: Opportunities for Predicting their Threat to Wildlife. *Conserv. Lett.*, 10: 477-484. doi:10.1111/conl.12297
- ROY, H.E., J. PEYTON, D.C. ALDRIDGE, T. BANTOCK, T.M. BLACKBURN, R. BRITTON, P. CLARK, E. COOK, K. DEHNEN-SCHMUTZ, T. DINES, M. DOBSON, F. EDWARDS, C. HARROWER, M.C. HARVEY, D. MINCHIN, D.G. NOBLE, D. PARROTT, M.J.O. POCKOCK, C.D. PRESTON, S. ROY, A. SALISBURY, K. SCHÖNRÖGGE, J. SEWELL, R.H. SHAW, P. STEBBING, A.J.A. STEWART & K.J. WALKER. 2014. Horizon scanning for invasive alien species with the potential to threaten biodiversity in Great Britain. *Global Change Biol.*, 20: 3859-3871.
- SAIZ, E., A. CALBET & E. BROGLIO. 2003. Effects of small-scale turbulence on copepods: The case of *Oithona davisae*. *Limnol. Oceanogr.*, 48(3): 1304-1311.
- SCHOFIELD, P.J. 2010. Update on geographic spread of invasive lionfishes (*Pterois volitans* [Linnaeus, 1758] and *P. miles* [Bennett, 1828]) in the Western North Atlantic Ocean, Caribbean Sea and Gulf of Mexico. *Aquat. Invasions*, 5: 117-122.
- SHIGANOVA, T., E. MUSAIEVA, T. LUKASHEVA, A.N. STUPNIKOVA, D.N. ZAS'KO, L.L. ANOKHINA, A.E. SIVKOVICH, V.I. GAGARIN & YU.V. BULGAKOVA. 2012. Increase in Findings of Mediterranean Non-native Species in the Black Sea. *Russ. J. Biol. Invasions*, 3: 255-280.
- SMITHSONIAN MARINE STATION AT FORT PIERCE. 2016. *Penaeus aztecus* (brown shrimp). http://www.sms.si.edu/irlspec/Penaeus_aztecus.htm (Accessed on 04/06/2016).

- STEIDINGER, K.A. & K. TANGEN. 1996. Dinoflagellates. In: C.R. Tomas (Editor). Identifying Marine Diatoms and Dinoflagellates. Academic Press, New York, pp. 387-598.
- STRASSER, M. 1999. *Mya arenaria* — an ancient invader of the North Sea coast. Helgol. Mar. Res., 52: 309-324.
- SVENSSON, J.R., G.M. NYLUND, G. CERVIN, G.B. TOTH & H. PAVIA. 2013. Novel chemical weapon of an exotic macroalga inhibits recruitment of native competitors in the invaded range. J. Ecol., 101(1): 140-148.
- SVISTUNOVA, L. 2013. A new invader in plankton of the Sea of Azov. Vestn. South. Centre, 9: 104-107.
- SYOMIN, V.L., A.V. SIKORSKI & A.I. SAVIKIN. 2017. *Streblospio* cf. *gynobranchiata* (Polychaeta, Spionidae) from the eastern part of Taganrog Bay (in Russian). Зоол. Ж./Zool. Zh., 96(1): 119-120.
- TAHERI, M., J. SEYFABADI, B. ABTAHI & M.Y. FOSH-TOMI. 2008. Population changes and reproduction of an alien spionid polychaete, *Streblospio gynobranchiata*, in shallow waters of the south Caspian Sea. Mar. Biodiv. Records, e40: 1-5.
- TAVARES, M. 2002. Shrimps. The Living Marine Resources of the Western Central Atlantic. In: K.E. Carpenter (Editor). Volume 1: Introduction. Molluscs, Crustaceans, Hagfishes, Sharks, Batoid Fishes and Chimaeras. FAO Species Identification Guide for Fisheries Purposes and American Society of Ichthyologists and Herpetologists Special Publication No. 5. Rome, Italy, FAO, pp. 251-291.
- TAYLOR, F.J.R., Y. FUKUYO & J. LARSEN. 1995. Taxonomy of harmful dinoflagellates. In: G.M. Hallegraeff, D.M. Anderson and A.D. Cembella (Editors). Manual on Harmful Marine Microalgae, IOC Manuals and Guides No. 33. UNESCO, France, pp. 283-317.
- TEACĂ, A., T. BEGUN & M. MURESAN. 2015. Chapter 7: Assessment of Soft-Bottom Communities and Ecological Quality Status Surrounding Constanta and Mangalia Ports (Black Sea). In: C. Stylios, T. Floqi, J. Marinski & L. Damiani (Editors). Sustainable Development of Sea-Corridors and Coastal Waters. Springer International Publishing, Switzerland, pp. 67-74.
- TRICHKOVA T., A. ULUDAĞ, A. ZENETOS, V. VLADIMIROV, R. TOMOV & D. COGALNICEANU (Editors). 2017. ESENIAS Scientific Reports 2. East and South European Network for Invasive Alien Species – A tool to support the management of alien species in Bulgaria (ESENIAS-TOOLS). IBER-BAS, ESENIAS, Sofia, Bulgaria, in press.
- ULMAN, A., J. FERRARIO, A. OCCHPINTI-AMBROGI, C. ARVANITIDIS, A. BANDI, M. BERTOLINO, C. BOGI, G. CHATZIGEORGIOU, B.A. ÇİÇEK, A. DEIDUN, A. RAMOS-ESPLÁ, C. KOÇAK, M. LORENTI, G. MARTINEZ-MAIZ, G. MERLO, E. PRINCISGH, G. SCRIBANO & A. MARCHINI. 2017. A massive update of non-indigenous species records in Mediterranean marinas. Peer J, 5: e3954.
- UYE, S. & K. SANO. 1995. Seasonal reproductive biology of the small cyclopoid copepod *Oithona davisae* in a temperate eutrophic inlet. Mar. Ecol. Progr. Ser., 118: 121-128.
- VELIKOVA, V., S. MONCHEVA & D. PETROVA. 1999. Phytoplankton dynamics and red tides (1987-1997) in the Bulgarian Black Sea. Water Sci. Technol., 39(8): 27-36.
- VERLAQUE, M., S. RUITTON, F. MINEUR & C.F. BOUDOURESQUE. 2015. Macrophytes. CIESM Atlas of exotic species in the Mediterranean. CIESM Publishers, Monaco, 364 pp.
- WALKER, L.M. & K.A. STEIDINGER. 1979. Sexual reproduction in the toxic dinoflagellate *Gonyaulax monilata*. J. Phycol. 15: 312-315.
- YOKES, B. & B. GALIL. 2006. The first record of the needlespined urchin *Diadema setosum* (Leske, 1778) (Echinodermata: Echinoidea: Diadematidae) from the Mediterranean Sea. Aquat. Invasions, 1(3): 188-190.
- ZENETOS, A. & P.K. KARACHLE. 2017. Marine non-indigenous species in South-East Europe. In: M. Rat, T. Trichkova, R. Scalera, R. Tomov and A. Uludag (Editors). ESENIAS Scientific Reports 1. State of the Art of Alien Species in South-East Europe. ESENIAS, pp. 123-136.
- ZENETOS, A., M. ARIANOUTSOU, I. BAZOS, S. BALOPOULOU, M. CORSINI-FOKA, M. DIMI-

- ZA, P. DRAKOPOULOU, S. KATSANEVAKIS, G. KONDYLATOS, N. KOUTSIKOS, E. KYTINOU, E. LEFKADITOU, M.A. PANCUCCI-PAPADOPOULOU, M. SALOMIDI, N. SIMBOURA, G. SKOUFAS, P. TRACHALAKIS, M. TRIANTAPHYLLOU, K. TSIAMIS, N.J. XENTIDIS & D. POURSANIDIS. 2015. ELNAIS: A collaborative network on Aquatic Alien Species in Hellas (Greece). *Manag. Biol. Invasion.*, (2): 185-196.
- ZENETOS, A., S. GOFAS, C. MORRI, A. ROSSO, D. VIOLANTI, J.E. GARCÍA RASO, M.E. ÇINAR, A. ALMOGI-LABIN, A.S. ATES, E. AZZURRO, E. BALLESTEROS, C.N. BIANCHI, M. BILECENOGU, M.C. GAMBI, A. GIANGRANDE, C. GRAVILLI, O. HYAMS-KAPHZAN, P.K. KARACHLE, S. KATSANEVAKIS, L. LIPEJ, F. MASTROTOTARO, F. MINEUR, M.A. PANCUCCI-PAPADOPOULOU, A. RAMOS ESPLÁ, C. SALAS, G. SAN MARTÍN, A. SFRISO, N. STREFTARIS & M. VERLAQUE. 2012. Alien species in the Mediterranean Sea by 2012. A contribution to the application of European Union's Marine Strategy Framework Directive (MSFD). Part 2. Introduction trends and pathways. *Medit. Mar. Sci.*, 13(2): 328-352.
- ZENETOS, A., S. GOFAS, M. VERLAQUE, M.E. CINAR, J.E. GARCÍA RASO, C.N. BIANCHI, C. MORRI, E. AZZURRO, M. BILECENOGU, C. FROGLIA, I. SIOKOU, D. VIOLANTI, A. SFRISO, G. SAN MARTÍN, A. GIANGRANDE, T. KATAĞAN, E. BALLESTEROS, A. RAMOS-ESPLÁ, F. MASTROTOTARO, O. OCAÑA, A. ZINGONE, M.C. GAMBI & N. STREFTARIS. 2010. Alien species in the Mediterranean Sea by 2010. A contribution to the application of European Union's Marine Strategy Framework Directive (MSFD). Part I. Spatial distribution. *Medit. Mar. Sci.*, 11(2): 381-493.
- ZIBROWIUS, H. 2002. Assessing scale and impact of ship transported alien fauna in the Mediterranean? CIESM Workshop Monographs, 20: 62-68.
- ZOLOTAREV, V. 1996. The Black Sea ecosystem changes related to the introduction of new mollusc species. *Mar. Ecol.*, 17: 227-236.

Uspostavljanje popisa morskih invazivnih vrsta u ESENIAS području: sadašnja situacija i buduća očekivanja

Paraskevi K. KARACHLE*, Maria CORSINI FOKA, Fabio CROCETTA,
Jakov DULČIĆ, Nina DZHEMBEKOVA, Marika GALANIDI, Petya IVANOVA,
Noa SHENKAR, Marius SKOLKA, Elitsa STEFANOVA, Kremena STEFANOVA, Victor
SURUGIU, Irfan UYSAL, Marc VERLAQUE i Argyro ZENETOS

**Kontakt e-adresa: pkarachle@hcmr.gr*

SAŽETAK

U ovom radu predstavljamo popis invazivnih i potencijalno invazivnih vrsta na istoku i jugu Europske mreže za invazivne vrste (ESENIAS) u zemljama s morskim granicama.

Vrste su klasificirane prema postojećoj literaturi i procjeni stručnjaka, pa su tako utvrđene grupe povremenih, invazivnih i očekivanih vrsta. Podaci su dati za deset vrsta čija se važnost temelji na njihovom širenju i invazivnom karakteru.

Od 160 vrsta koje sadrži popis, 148 je već bilo prisutno u ESENIAS zemljama, dok je 9 invazivnih vrsta bilo prisutno u Sredozemnom ili u drugim europskim morima, a vjerojatno je da će se zabilježiti i u zemljama udruženim u ESENIAS. Većina vrsta je bile iz Crvenog mora / indopacifičkog podrijetla (97 vrsta, 60,6%). Italija, Turska i Grčka su zemlje s najvišom zastupljenošću vrsta (159, 152 i 139), zbog njihove proširene obale i broja znanstvenika koji rade na morskim invazivnim vrstama.

Najveći broj utvrđenih vrsta zabilježen je u Turskoj (116 vrsta), dok su u Italiji i Grčkoj najbrojnije vrste bile "očekivane" (85 i 48 vrsta, respektivno).

Istočne zemlje Jadranskog mora (npr. Albanija, Hrvatska, Crna Gora i Slovenija) imale su općenito nizak broj vrsta na ovom popisu, od kojih mnoge još "očekujemo" da pristignu iz susjednih zemalja: Grčke i Italije.

Konačno, najčešće su potencijalni putovi bili „transforni putnici“ (balastna voda broda: 41 slučaj, obraštaj brodskog trupa: 55), dok je slijedilo i širenje lesepsijskih migranata (95 slučajeva).

Ovaj popis je namijenjen da služi kao sustav ranog upozorenja koji bi kroz proces skeniranja pomogao državama ESENIAS da daju prioritet invazivnim stranim vrstama, njihovim putovima i područjima veće vjerojatnosti pojavljivanja, kako bi se poduzele potrebne mjere upravljanja.

Ključne riječi: invazivne vrste, ESENIAS, Sredozemno more, Crno more

