



Monitoring Extreme Impacts of *Rugulopteryx okamurae* (Dictyotales, Ochrophyta) in El Estrecho Natural Park (Biosphere Reserve). Showing Radical Changes in the Underwater Seascape

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Specialty section:

This article was submitted to
Biogeography and Macroecology,
a section of the journal
Frontiers in Ecology and Evolution

Received: 08 December 2020

Accepted: 12 March 2021

Published: 15 April 2021

Citation:

García-Gómez JC, Florido M, Olaya-Ponzzone L, Rey Díaz de Rada J, Donázar-Aramendía I, Chacón M, Quintero JJ, Magariño S and Megina C (2021) Monitoring Extreme Impacts of *Rugulopteryx okamurae* (Dictyotales, Ochrophyta) in El Estrecho Natural Park (Biosphere Reserve). Showing Radical Changes in the Underwater Seascape. *Front. Ecol. Evol.* 9:639161. doi: 10.3389/fevo.2021.639161

The invasive macroalga *Rugulopteryx okamurae* represents an unprecedented case of bioinvasion by marine macroalgae facing the European coasts. Since the first apparition of the species in the Strait of Gibraltar in 2015, its fast dispersion along the introduced habitats constitutes a real challenge to develop monitoring strategies that ahead of its impacts. The present study uses three different approaches to address impacts on the benthic ecosystems, at the same time offers relevant data for future management actions in El Estrecho Natural Park (PNE). Information obtained by monitoring permanent sentinel stations revealed a significant loss in resident species coverage after the moment of maximum growth in 2017. Thus, despite coverage of *R. okamurae* did not strongly varied in the latter years, impacts generated remain high in the habitats studied. Estimations of the invasive species coverage by combining cartographic image analysis and *in situ* data predicted a major occupation (over 85% coverage) between 10 and 30 m, coinciding with the maximum rocky surface areas (m²) mapped on the PNE. Furthermore, a Citizen Science research collaboration evidenced impacts on the benthic seascape through an *ad hoc* exploration of images that allowed a “before” and “after” comparison of the invasion process in the same geographic locations. This has made it possible to graphically demonstrate severe changes in the underwater seascape and, therefore, the general impact of this new biological invasion. The spatial colonization estimations combined with the impacts reported by both scientific [Sessile Bioindicators in Permanent Quadrats (SBPQ) sentinel stations] and civilian (Citizen Science) monitoring methodologies claim the urgent development of further studies that allow the design of monitoring strategies against *R. okamurae* expansion across the Mediterranean and Atlantic waters.

Keywords: *Rugulopteryx okamurae*, Citizen Science, benthic biota, monitoring, invasive species, invasive macroalgae, El Estrecho Natural Park, Strait of Gibraltar

INTRODUCTION

The Strait of Gibraltar is a hot spot area for marine biodiversity in the Atlantic-Mediterranean waters with great biogeographic importance due to the coalescence of the Lusitanian, Mediterranean and Mauritanian regions (Ekman, 1953; Briggs, 1974; García-Gómez, 2002). Together with the Alboran Sea, the Strait harbors species from the northern Atlantic, subtropical waters of the northwestern of Africa and the Mediterranean Sea (Templado, 2011; Mannino et al., 2017; Urrea et al., 2017). Its species richness and habitat diversity is influenced by the littoral physiographic complexity, which has led to the co-occurrence and dominance of different biological strategies and thus affecting the composition of benthic communities (Zabala and Ballesteros, 1989). The biodiversity of this area has been largely assessed by a list of references in terms of foundation species [macroalgae (Flores-Moya et al., 1995a,b; Conde et al., 1996; Adama et al., 2021) and sessile macroinvertebrates as ascidians (Carballo et al., 1997) or molluscs (García-Gómez, 2002; Gofas et al., 2011)] and associated fauna (e.g., García-Raso, 1993; Conradi et al., 2000; Castello and Carballo, 2001; Guerra-García et al., 2009; García-Raso et al., 2011).

However, the benthic systems have changed over time according to different biotic and abiotic factors (see Gallardo et al., 2021). The biodiversity and species richness have been threatened by several anthropogenic pressures for many years, mainly due to the proximity to pollution resources (e.g., chemical industries, thermal plants, oil slick) (Morales, 2007; Soussi et al., 2020), artificial infrastructures (Sánchez-Moyano et al., 2000; Sedano et al., 2020) or maritime shipping and human activities (Bianchi et al., 2013; Nachite et al., 2020). Indeed, the Strait of Gibraltar supports the highest density of maritime traffic in the Western Mediterranean (Abdulla and Linden, 2008; Endrina et al., 2018). This implies a threat due to marine bioinvasions since ballast waters and boat hulls fouling are recognized as potential transport vectors for non-indigenous species (NIS) (Ribera-Siguan, 2003).

Favorable environmental conditions increase the settlement success of invaders (Villèle and Verlaque, 1995), which in turn compromise the survival of surface-dependent organisms by changing the physical characteristics and available substrata (Mannino et al., 2017). Macrophytes (macroalgae and seagrasses) constitute the dominant group of invasive species in the Western Mediterranean (Zenetos et al., 2010, 2012). In the recent years, a number of studies have been performed on invasive macroalgae that have interacted the resident sessile communities promoting also cascading influence on associated fauna in the Strait waters. Some examples are *Asparagopsis armata* Harvey (Boudouresque and Verlaque, 2002; Guerra-García et al., 2012), *Asparagopsis taxiformis* (Delile) Trevisan de Saint-Léon (Altamirano et al., 2008; Navarro-Barranco et al., 2018), *Caulerpa cylindracea* Sonder (Altamirano et al., 2014); *Womersleyella setacea* (Hollenberg) R. E. Norris (Bedini et al., 2015), or *Lophocladia lallemandii* (Montagne) F. Schmitz (Ballesteros et al., 2007). Despite positive and neutral effects may also occur [e.g., *Caulerpa racemosa* (Forsskål) J. Agardh may maintain caprellid populations in shallow Mediterranean habitats (Vázquez-Luis

et al., 2009)], these species have displayed pronounced and drastic effects on the underwater seascape making it difficult to implement success of strategies to mitigate their impacts (Anderson, 2007; Klein and Verlaque, 2008).

The last notorious case of macroalga colonizing the Western Mediterranean littorals is the species *Rugulopteryx okamuræ* (E. Y. Dawson) I.K. Hwang, W.J. Lee, and H.S. Kim. This brown macroalga, originated from the northwest of Asia (Hwang et al., 2009), has been systematically reported on the coasts of the Strait of Gibraltar since 2015 (see Altamirano-Jeschke et al., 2016), when more than 5,000 tons of wracks were extracted from the beaches of its south coasts, in the city of Ceuta (North-Africa) (Ocaña et al., 2016). The rocky bottoms of The Jbel Moussa Site of Biological and Ecological Interest (SIBE) (El Aamri et al., 2018) and the eastern littoral of El Estrecho Natural Park (PNE) (García-Gómez et al., 2018) firstly represented the northern and southern scenarios of *R. okamuræ* expansion in the Strait of Gibraltar. Until date, these areas constitute the most intensely affected by the brown alga, which continues its westward and eastward directionality of expansion (Altamirano et al., 2019; Figueroa et al., 2020) with trend to monopolize the sea rocky bottom in detriment of the photophilous resident biota (García-Gómez et al., 2020b). In the introduced areas, the species is present throughout the year and is dispersed mainly due to asexual and vegetative strategies by propagules and monospores (Altamirano-Jeschke et al., 2017; Altamirano et al., 2019). Although it has not been possible to assess if the species is able to complete its entire life cycle in the Atlantic and Mediterranean waters (Verlaque et al., 2009; Altamirano-Jeschke et al., 2016, 2017), the fast expansion and massive occupation potential since its first detection in 2015 reflects that this bioinvasion case is one of the most serious and threatening caused by marine macroalgae in the European waters (García-Gómez et al., 2018, 2020b).

It is urgent to carry out studies on the distribution, ecology and impacts of *R. okamuræ* in the Mediterranean and the Atlantic coasts, as well as the implementation of management measures. However, few studies have been carried out on *R. okamuræ* distribution (e.g., Altamirano-Jeschke et al., 2016; Ocaña et al., 2016; El Aamri et al., 2018; Altamirano et al., 2019) and its derived impacts on the recipient sessile (García-Gómez et al., 2018, 2020b; Sempere-Valverde et al., 2020) and mobile associated biota (Navarro-Barranco et al., 2019). In this regard, there is only one published study monitoring the temporal dynamic of the invasion since the first apparition of the species in the Strait waters (see García-Gómez et al., 2020b) by the utilization of Sessile Bioindicators in Permanent Quadrats (SBPQ). The SBPQ is part of a monitoring of sessile sentinel species which has been carried out since 2013 to detect early impacts on the littoral environment, including changes in the benthic system related to global warming (García-Gómez, 2015). Thus, this methodology has allowed to document not only the establishment of *R. okamuræ* but also the disappearance of target species which become displaced in the absence of environmental stability.

Long-term monitorings, as observational studies that obtain full ecological characterizations in environmental evaluations

(Moschella et al., 2005), are essential for the management of invasive species, their early detection and the implementation of a rapid response (Lodge et al., 2006; Williams and Smith, 2007; García-Gómez, 2015). In this sense, Citizen Science is a research collaboration strategy involving members of the public in scientific research projects to address real-world problems (Wiggins and Crowston, 2011). This constitutes an emerging movement of citizen participation given the breakdown or reduction of barriers caused by advances in communication technologies. Although there are challenges to effectively use citizen-generated data to monitor invasive species on a global scale (Earp and Liconti, 2020; Johnson et al., 2020), it can be a useful tool to detect and monitoring the bioinvasion processes (present study). In this sense, special emphasis has been placed on the underwater seascape, as highlighted in previous works performed in other research fields (Pittman et al., 2011; Gobert et al., 2014; Cheminée et al., 2016; Schejter et al., 2017; Ceraulo et al., 2018). The monitoring of an area by comparative images reflecting “before” and “after” scenarios due to bioinvasion processes as a proxy of BACI analyses (Before After Control Impact) (Underwood, 1994; Montefalcone et al., 2008; Conner et al., 2015; Donázar-Aramendía et al., 2018, 2020), may offer key information on the degree of affection that the local ecosystem has suffered and the risk of its prevalence in the area. Among other aspects, the comparative analysis of images can allow inferences regarding the behavior of biota and ecological connectivity, since the underwater seascape can have a great influence on it (Grober-Dunsmore et al., 2009).

The overall aim of this study was to advance our knowledge on the *R. okamuræ* bioinvasion case targeting efforts on different monitoring approaches. Thus, objectives were threefold: (1) to provide updated data about the evolution of *R. okamuræ* and key resident taxa in pre-coralligenous habitats and to test better designs for future monitoring programs; (2) to estimate the coverage of the invasive species on the rocky surface areas of the PNE, providing mapping areas of suitable rocky substrata to host the invasive species; and (3) to offer graphical evidences of radical changes before and after the spatial establishment of *R. okamuræ* along the PNE littoral through Citizen Science collaborations.

MATERIALS AND METHODS

Study Area

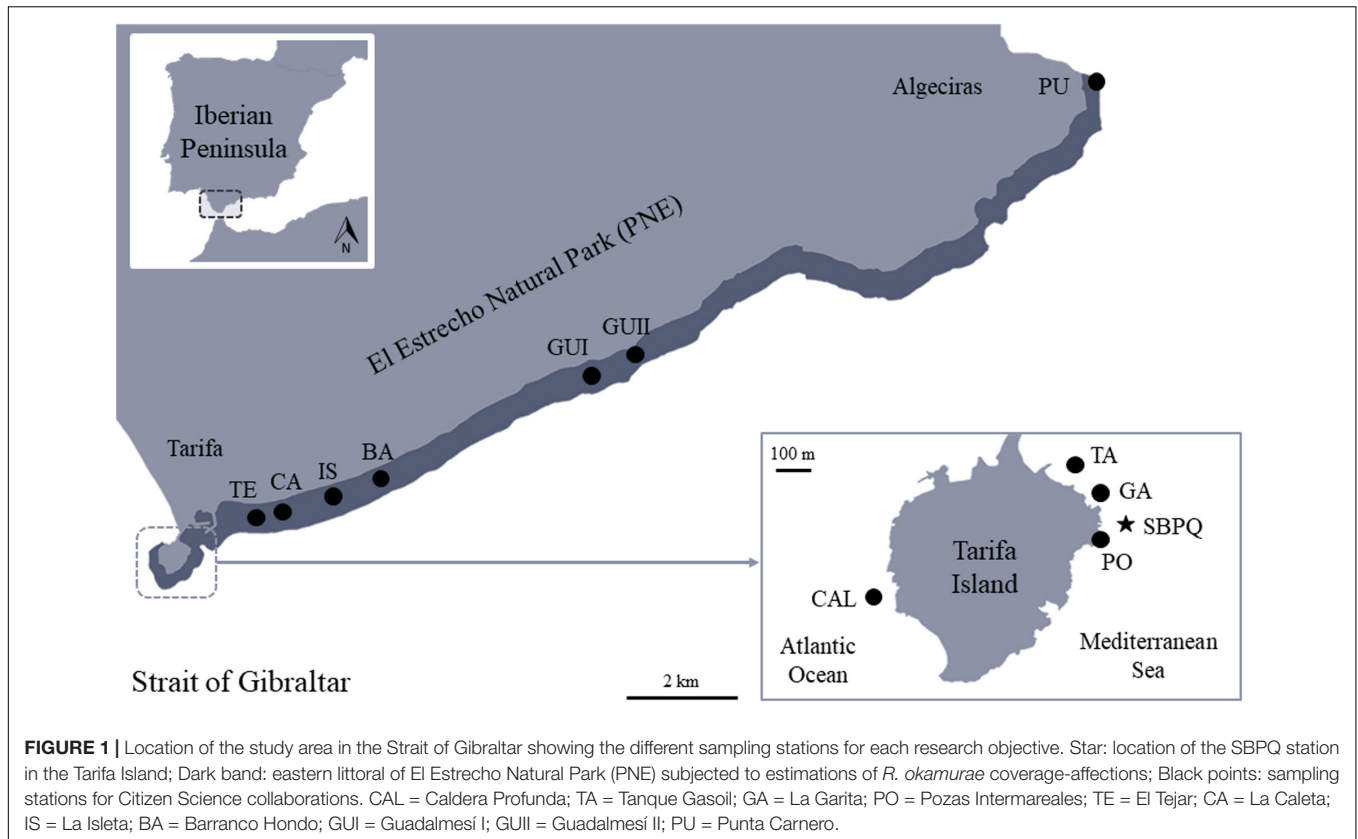
The present study is focused on the northern coastline of the Strait of Gibraltar. Concretely, the PNE, which is located in the southern Iberian Peninsula and included within the Marine Protected Areas (MPAs) of the Andalusian Intercontinental Biosphere Reserve (Spain-Morocco). Situated between the Atlantic Ocean (western coasts) and the Mediterranean Sea (eastern coasts), the PNE extends from southern Spain to northern Morocco, separating Europe and Africa by 14 km distance. The study area includes the eastern littoral of the PNE coastline, between Tarifa Island (36°05′20.38″N, 5°48′45.34″O) and Punta Carnero (Algeciras) (36°4′30.18″N, 5°25′11.16″O) (Figure 1).

Monitoring of the Submarine SBPQ Station of Tarifa Island

The sessile community at the SBPQ station of Tarifa Island (Figure 1) was seasonally sampled on partially shaded vertical walls of the pre-coralligenous habitats (15 m depth) during 2013–2017 by García-Gómez et al. (2020b). The SBPQ methodology is specifically designed to assess impacts on sensitive species characterized by long-life cycles, thus providing evidences in any period of the year. This methodology has been designed as a simple, non-invasive, underwater environmental alert tool for the potential early detection of environmental impacts of anthropic origin in the sublittoral system: in the short term (local alterations derived from pollutants, coastal dredging or civil engineering works, early detection of exotic species with invasive potential, etc.), and in the medium or long term (global warming) (García-Gómez, 2015; García-Gómez et al., 2020a). Because *R. okamuræ* is present all-year round in the Strait of Gibraltar (pers. obs.) and most of target species from the sentinel station of Tarifa showed at least 10% of coverage in all seasons (see García-Gómez et al., 2020b), it could be expected that this long-term technique based on permanent quadrats will allow the detection of changes in the coverage of the invasive and the target species only considering one sampling season. In this study, the monitoring activities started in 2013 were temporally continued until 2020 and only considering winters (no data are available from 2018, when sampling was missed because of technical issues).

The monitoring station of Tarifa Island is composed of three 1 × 1 m fixed quadrats separated 5–10 m from each other. Within each fixed quadrat, four photo-quadrats of 50 × 50 cm were collected using *in situ* photographs via scuba diving. Following the methodology proposed by García-Gómez (2015), García-Gómez et al. (2020a) and the monitoring study performed for 2013–2017 (García-Gómez et al., 2020b), species coverage (percent) was obtained by superposing 10 × 10 grid over each photo-quadrat obtained for each fixed quadrat. Since the methodology overestimates the species-cover, the total percentage coverage in each photo-quadrat exceeded 100%.

To visualize the spatial distribution of replicates (quadrats) among years, a non-metric multidimensional scaling (nmMDS) based on a Bray–Curtis similarity matrix was performed on the square root transformed coverage data considering all the data pool since 2013. Differences in the sessile community structure over the sampling years were analyzed using PERMANOVA analysis under a nested design (sampling times were defined as random factor, “Season,” nested within “Year,” fixed factor with seven levels: 2013–2020). P-values were obtained through a Monte Carlo test when small number of unique permutations were obtained (Anderson et al., 2008). Also, the homogeneity of the data dispersion among samples was tested for the fixed factor “Year,” using a permutational analysis of multivariate dispersions (PERMDISP) (Anderson, 2006). The contribution of each species to the Bray–Curtis similarity was obtained using a SIMPER (SiMilarity PERcentages) analysis. After performing analyses considering the whole data base, we wanted to test if the methodology was functional only sampling once a year considering the community present in this sentinel station. We



considered winters as the season of the year most appropriate to test whether the SBPQ methodology is minimally robust, since it is the time when it is expected (for middle latitudes) that the development and growth of the invasive macroalga is lower (this has been supported by pers. obs. and growth data for *R. okamuræ* in Japan (Agatsuma et al., 2005), with similar north latitude to Spain: between 20° and 40°). Thus, the sensitivity of the SBPQ methodology was tested repeating all the analyses but only considering the data from the winter of each year (i.e., factor “Year,” with seven levels: 2013–2020). Overall multivariate analyses were carried out using PRIMER-e v6 PERMANOVA + software (Clarke and Gorley, 2006; Anderson et al., 2008).

Estimations of *Rugulopteryx okamuræ* Coverage in the PNE

The sublittoral physical cartography of the Strait of Gibraltar (0–1 nautical mile zone) (García-Gómez et al., 2003; see also CMAyOT Consejería de Medio Ambiente y Ordenación del Territorio, 2008; MITECO Ministerio para la Transición Ecológica, 2012) was used to estimate the geographical and bathymetric expansion of *R. okamuræ* along the sublittoral coastline of the PNE (Figure 1). Thus, *R. okamuræ* coverage data were estimated and applied to bathymetric intervals of rocky sublittoral habitats (0–5, 5–10, 10–20, 20–30, and 30–40 m) by averaging coverage data obtained at the horizontal illuminated rocky surfaces of the southern coasts of the Strait (summer 2018, at 14 km from the study area; data previously published in García-Gómez et al., 2020b). Thus, the

degree of coverage of the rocky bottoms were mapped using a polygon methodology for each bathymetric interval considered as follow: (1) A digital bathymetry model (DTM) was made in the PNE area and a 3-m orthogonal mesh was generated (surfer application from Golden Software, LLC was used). (2) The polygons of the outcropping rocks were extracted from the GIS database and (3) an intersection operation was performed between the DTM and the outcropping rock polygons. For this, the geo-processing tools of the Qgis application with the GNU general public license was used. (4) Once the DTM of the bathymetry was obtained, surface of the outcropping rock in each bathymetric interval was automatically calculated using a basic scripting language code from Golden Software, LLC (Contarea2.bas, 2000). The planar area (projected surface in the horizontal plane) and the real surface were calculated. (5). Finally, to generate a mapping of the rocky areas, the bathymetric DTM was divided in the ranges described above, so percentage coverages per bathymetric interval considered were displayed.

“Before-After” Underwater Seascape Approximation by Citizen Science Initiative

With the aid of local scuba divers and citizen scientists, the research group of this study often obtains useful information in environmental assessments on the Strait coasts and Algeciras bay. In this regard, occasional observational surveys of the benthic communities have been performing on the subtidal coastline of the PNE for more than 10 years, providing valuable

evidences of the underwater seascape dynamic over time. In this study, 11 video graphically monitored sites between Tarifa and Punta Carnero (Algeciras) (**Figure 1**) have been selected to compare environmental scenarios before and after the first establishment of *R. okamuræ* on the northern Strait of Gibraltar (2015). Citizen participants were constituted by experienced local divers, that carefully recorded the underwater seascape every time they performed their activities in the sites indicated. With the attempt to combine minimal diving efforts with the widest relevant information obtained (and thus, an optimal collaboration), data was recorded by video graphic instead of photographic methods (Cabatain et al., 2006). Thus, citizen and scientific collaborators provided a large number of video graphic data, of which individual video frames were selected and analyzed in the laboratory.

Data from videos recorded between Tarifa Island and Punta Carnero (i.e., El Tejar, La Caleta, La Isleta, Barranco Hondo, Guadalmesí I, Guadalmesí II and Punta Carnero) yielded information on the whole natural rocky area between 3 and 10 m depth, while videos from Tarifa Island offered data on artificial substrata (i.e., Tanque Gasoil and Caldera Profunda; 7–10 m depth) and natural sublittoral (i.e., La Garita; 10–14 m depth) and intertidal habitats (i.e., Pozas Intermareales). Images taken before and after the first citation of *R. okamuræ* in the Strait coasts (2015) were then selected and compared providing different scenarios of the underwater seascape and bioinvasion consequences.

RESULTS

Sessile Communities at the SBPQ Monitoring Station of Tarifa (2013–2020)

Results obtained showed that the percentage coverage of the sessile species at the SBPQ sentinel station located on vertical surfaces of shady pre-coralligenous habitats in Tarifa Island suffered a decrement in terms of species percent coverage until 2020, excepting *R. okamuræ* and the coral *Astroides calycularis* (Pallas, 1766), which slightly increased after 2019 (**Figure 2**). PERMANOVA results revealed no evidences of significant differences in the community structure between 2019 and 2020. However, it significantly differed from previous years (2013–2017) (**Table 1A** and **Figure 3A**). The same pattern was observed when the community from 2017 was compared with the rest of sampling years. SIMPER analysis revealed that dissimilarities with previous years were mainly given by *R. okamuræ* mean coverage values, which highly increased until 2017 (**Supplementary Table 1**). Instead, *R. okamuræ* coverage decreased after 2017, and only contributed to differences with most recent data (2019–2020) in 13 and 15%, respectively. In this case, it was the native community which mainly contributed to dissimilarities, with an accumulative contribution up to 60% in both cases and lower average values.

Results obtained only considering samplings from winters are shown in **Supplementary Figure 1** and **Table 1B**. Comparisons revealed a hindrance of the exponential growth of *R. okamuræ* from 2016 (**Supplementary Figure 1**). In 2019 and 2020, a slightly coverage enhancement was observed, but these values

only contributed to dissimilarities against previous winters in a 16% (2019) and 17% (2020) (**Supplementary Table 2**). Overall, species coverage did not significantly differ between 2019 and 2020 (**Table 1B**). In fact, ordination analyses (nmMDS) revealed similar patterns to those obtained when including all the seasons from the period 2013–2017 (**Figure 3B**). Communities in 2020 significantly differed from winters before 2017 (included) and 2019 significantly differed from winters before 2016 (included). Despite the high coverages reached in 2019 and 2020 contributed to main differences against 2016 in both cases, when comparing with 2017, the loss of autochthonous brown algae and *A. calycularis* contributed up to 50% of dissimilarities in both cases.

Rugulopteryx okamuræ Coverage Estimations on the Sublittoral Areas of the PNE

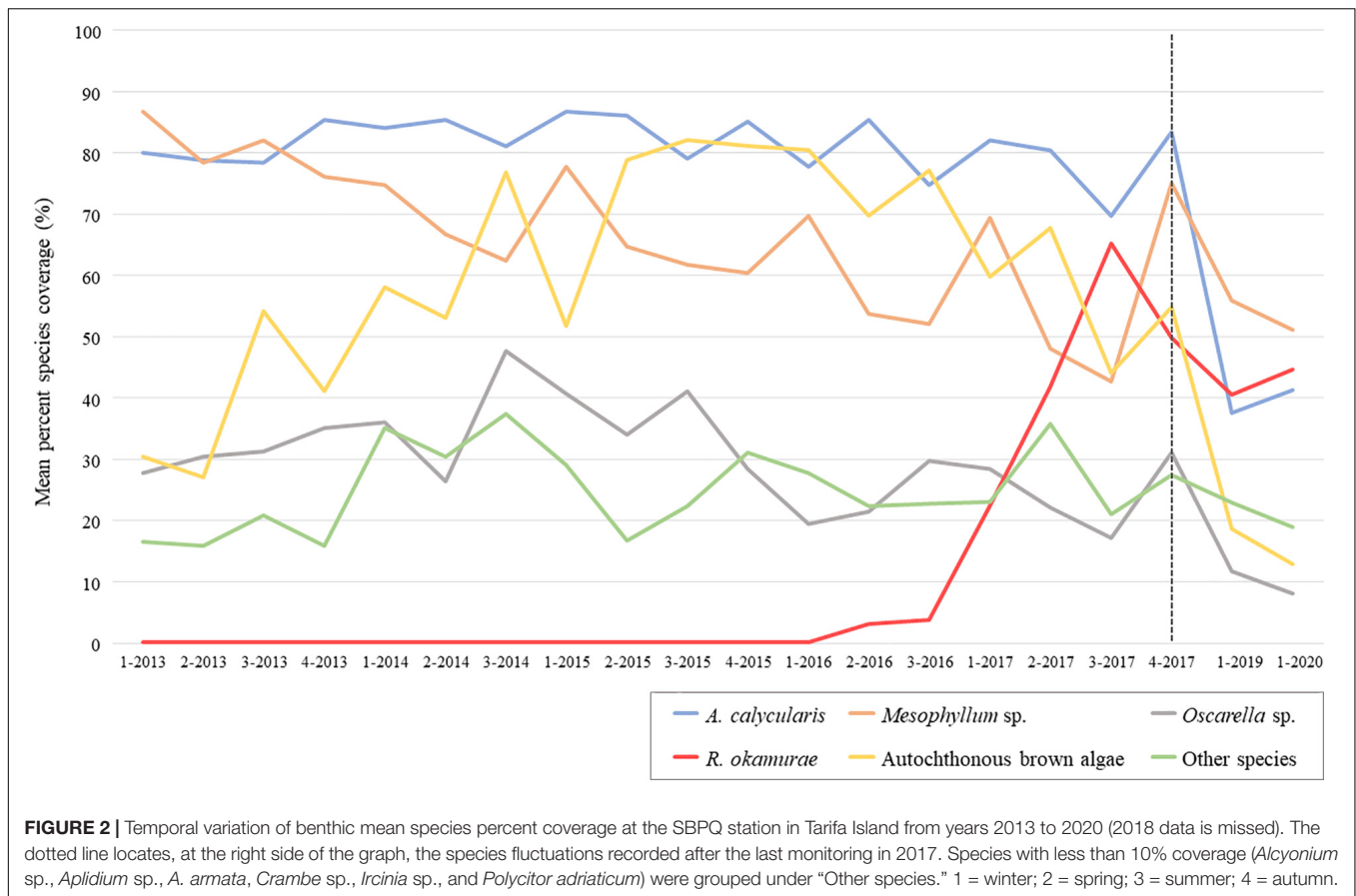
Coverage values registered at the bathymetric range of distribution of *R. okamuræ* in 2018 on horizontal illuminated surfaces (García-Gómez et al., 2020b) (**Figure 4A**) allowed the estimation of the species coverage for the different bathymetric intervals (**Figure 4B**). *R. okamuræ* percentage coverages were mapped for all the PNE coastline (**Figure 5**) and rocky surface areas and corresponding estimated percent coverages were obtained for the eastern littoral of the PNE (**Figures 6, 7**). The highest percentage coverages (85–96%) were obtained between 5 and 30 m depth. Over 10 m, these values coincided with the less extensive (565,469 m² between 5 and 10 m) and the most extensive rocky surfaces registered along the PNE (more than 3,000,000 m²) (**Figure 7**). Thus, 96% of total coverage values were reached at the bathymetric interval of 10–20 m depth, estimated for a total rocky surface of 3,141,476 m². Estimated percentages of *R. okamuræ* cover remained high (85%) until – 30 m. The lowest percentage coverages (42 and 45%) occupied the deepest and the shallowest habitats (30–40 m and 0–5 m, respectively). No coverage data deeper than 40 m were obtained since no measurements were performed at such depth in studies performed in 2018 (see García-Gómez et al., 2020b).

“Before-After” Underwater Seascape Approximation by Citizen Science Initiatives

The monitoring activities performed under the Citizen Science approach led to identify radical changes in the underwater seascape along the PNE littoral. Graphical data before and after the first establishment of *R. okamuræ* revealed visual changes in the benthic community composition where the invasive species was represented as a dominant component by most of image data taken after 2015. The selection of visual data and concerning results at each sampling station and time is presented in the **Table 2**.

DISCUSSION

Results obtained by the different objectives prospecting in this study evidenced the extreme effects of *R. okamuræ* on the benthic



ecosystems of the northern coasts of the Strait of Gibraltar, both by scientific (SBPQ alert method) and citizen (Citizen Science) monitoring methodologies. This, together with the high coverage levels estimated along the PNE coastline, illustrates the potential risk that the species represents to the benthic ecosystems in the Strait waters.

Updated Fluctuations of *Rugulopteryx okamurae* by Monitoring Sessile Bioindicators in Permanent Quadrats

The application of the SBPQ methodology has allowed the continuity of monitoring activities from 2017 to 2020 at the shady vertical surfaces of the pre-coralligenous habitats from the submarine sentinel station of Tarifa Island. Results showed that the benthic community structure differed between 2016 and 2017 due the high coverage levels of *R. okamurae* reached in that year. This pattern reflects the high spatial growth of the invasive species and, potentially, the ability to efficient using resources in the recipient habitats (Vaz-Pinto et al., 2014), which could results in the stronger and faster growth in detriment of the resident community, as is observed later in 2019 and 2020.

Until our study, few data have been offered about dominance-dynamics of *R. okamurae* after the huge coverages reached in 2017. Using the same methodology, Sempere-Valverde et al.

(2020) registered coverage increments of the invasive species in most of monitored sites of coralligenous habitats after 2017. Our results showed a decrease in percent coverage, slightly increasing in 2020. Changes in shape and size of the macroalgal beds can be modified by a variety of abiotic factors as temperature, salinity or turbidity (Glasby et al., 2005a). In fact, according to bloom-bust dynamic theories, a drastic decline in the invasive populations can occur after an initial rapid increase of the abundances, without implying recoveries in the macroalgal resident communities [e.g., *Caulerpa taxifolia* (M.Vahl) C.Agardh (Glasby et al., 2005a,b)]. This has been further combined with other works that assume that impacts on the resident biota cannot be assessed only considering species gain and loss without including mechanisms involved (Chapin et al., 2000). According to our results, differences in the community structure between the last 2 years and 2017 were due to an effective loss of resident biota, instead of fluctuations in *R. okamurae* presence. This is in line with results from Sempere-Valverde et al. (2020), who also found changes in the community structure and the regression of bioindicator species. Thus, despite certain coverage stabilization could have been raised in the last years, impacts generated by *R. okamurae* remains high in the habitat studied, and therefore it can be assumed that no signs of decline in its invasive potential have been perceived. However, longer-scale monitorings are needed in order to totally establish the strength of the bioinvasion, so

TABLE 1 | PERMANOVA and PERMDISP (for the factor “Year”) results for SBPQ station coverage data (2013–2020) **(A)** considering intra-annual variability (i.e., all seasons) and **(B)** only winter seasons.

	df	SS	MS	Pseudo-F	P(MC)
(A)					
Year	6	5,329.8	888.3	18.673	<0.001
Time (year)	13	618.43	47.571	0.97385	0.5173
Res	40	1,954	48.849		
Total	59	7,902.2			
P (PERMDISP):	0.6006				
Pair-wise test for years 2019 and 2020	2019 = 2020 ≠ 2017 ≠ (2016, 2015, 2014, 2013)				
	2020 = 2019 ≠ 2017 ≠ (2016, 2015, 2014, 2013)				
(B)					
Year	6	2,995.1	499.18	8.2181	<0.001
Res	14	850.38	60.741		
Total	20	3,845.5			
P (PERMDISP)	0.4237				
Pair-wise test for years 2019 and 2020	2019 = 2020; = 2017; ≠ (2016, 2015, 2014, 2013)				
	2020 = 2019; ≠ 2017; ≠ (2016, 2015, 2014, 2013)				

Pair-wise comparisons are based in Monte-Carlo P values. Summary results of Pair-Wise tests are showed only for comparisons between 2019–2020 and the series 2013–2017 (evidences of significant differences are referred as ≠ [i.e., P(MC) < 0.05] while “=” is used when no evidences were obtained).

monitoring efforts should be continued in the future to full characterize dynamics observed here.

Sessile Bioindicators in Permanent Quadrats focuses on monitoring information in substrata colonized by colonial species with long life cycle [i.e., pre-coralligenous (García-Gómez et al., 2020a) and coralligenous rocky bottoms (Sempere-Valverde et al., 2020)], so changes due to long established species can be easily perceived. In this study, the data available in winter revealed similar patterns while more progressive changes across years than when considering the whole intra-annual variability from 2013 to 2017. Thus, although the larger information obtained the higher characterization of the species dynamic can be ensured, the high similarity between ordination analyses performed in this study, reinforce the utility of the SBPQ methodology even with one sampling per year. Moreover, taking into account that at least in introduced habitats the species is also present in winter, cold seasons could be especially interesting because smaller coverages of the species may allow to reflect competitive processes missed in those periods where huge biomass led the substrata overgrowth.

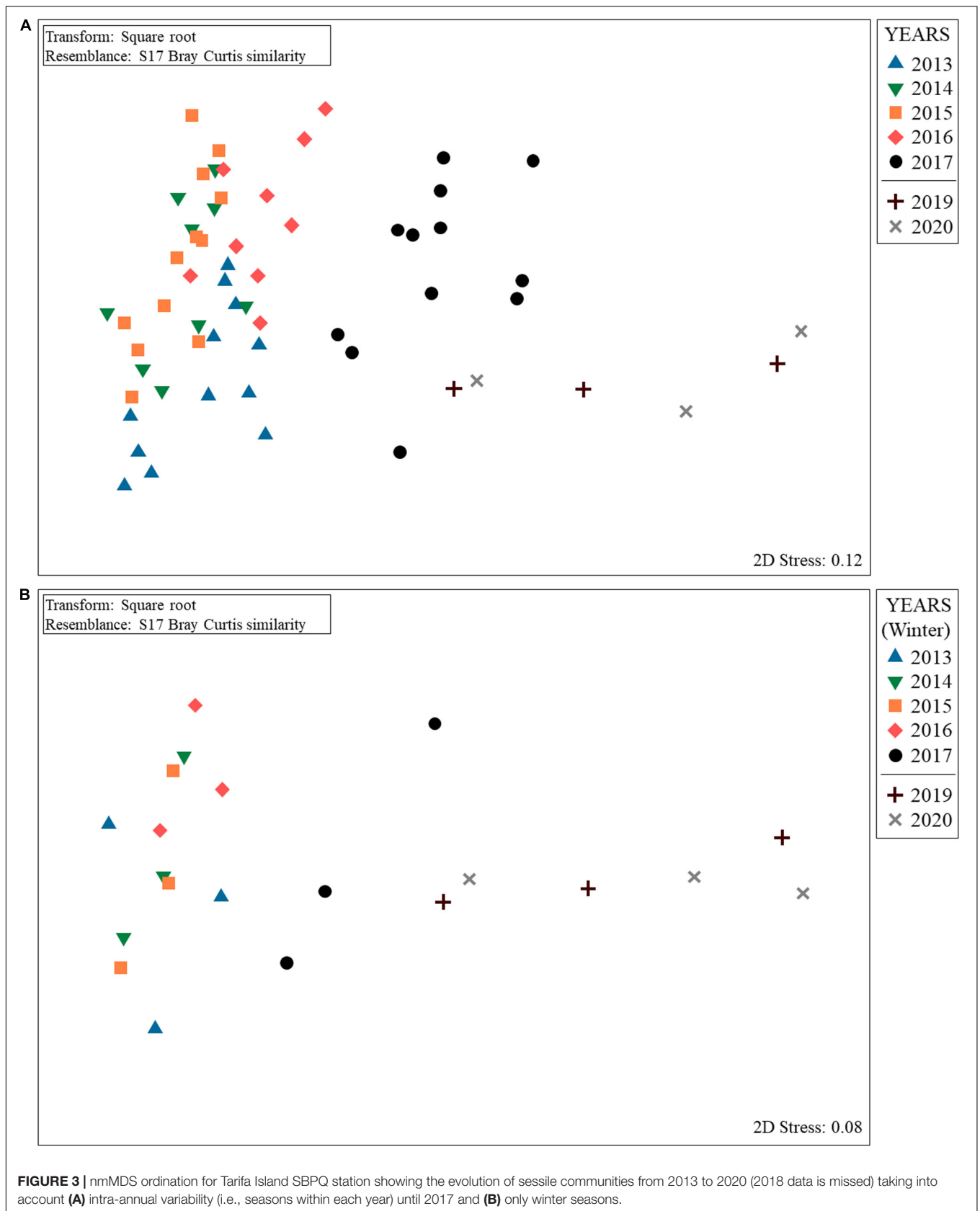
Implications of Coverage Estimations of *Rugulopteryx okamuræ* Within the PNE

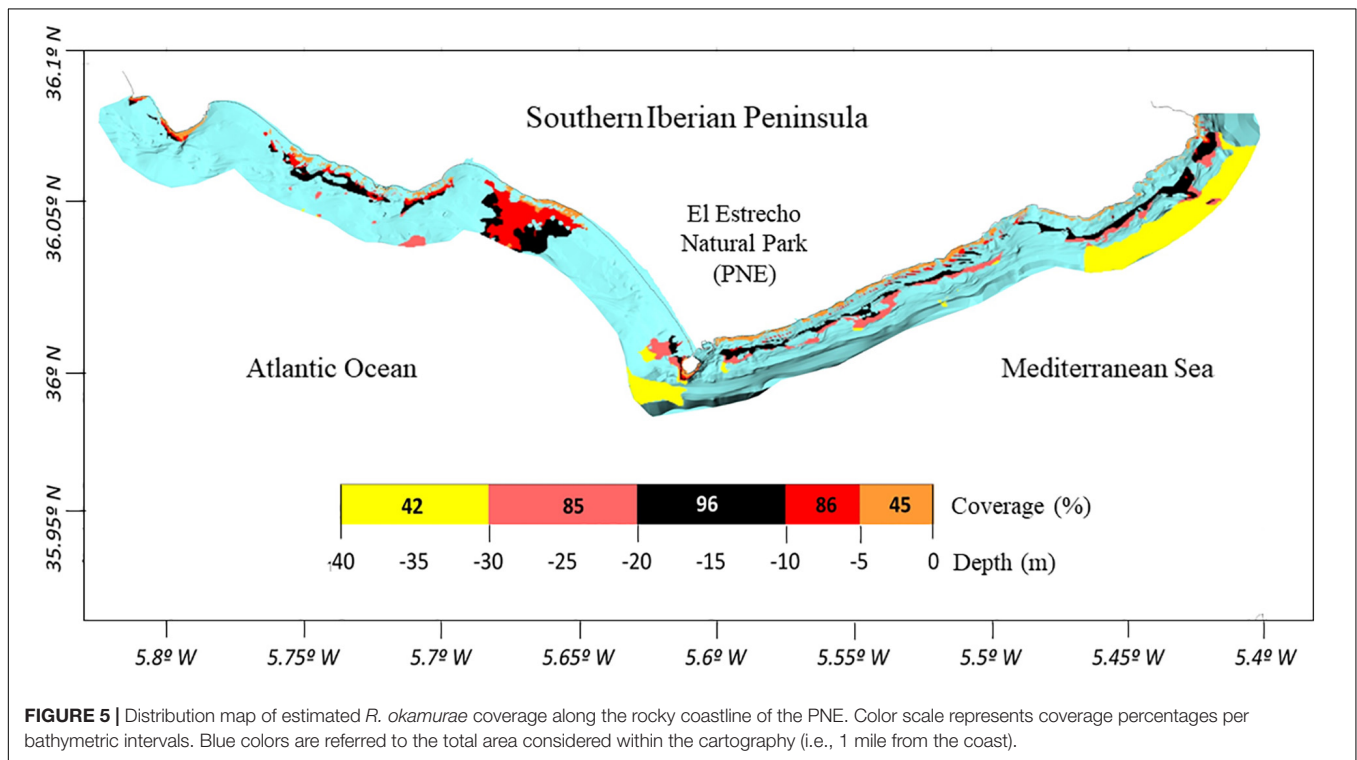
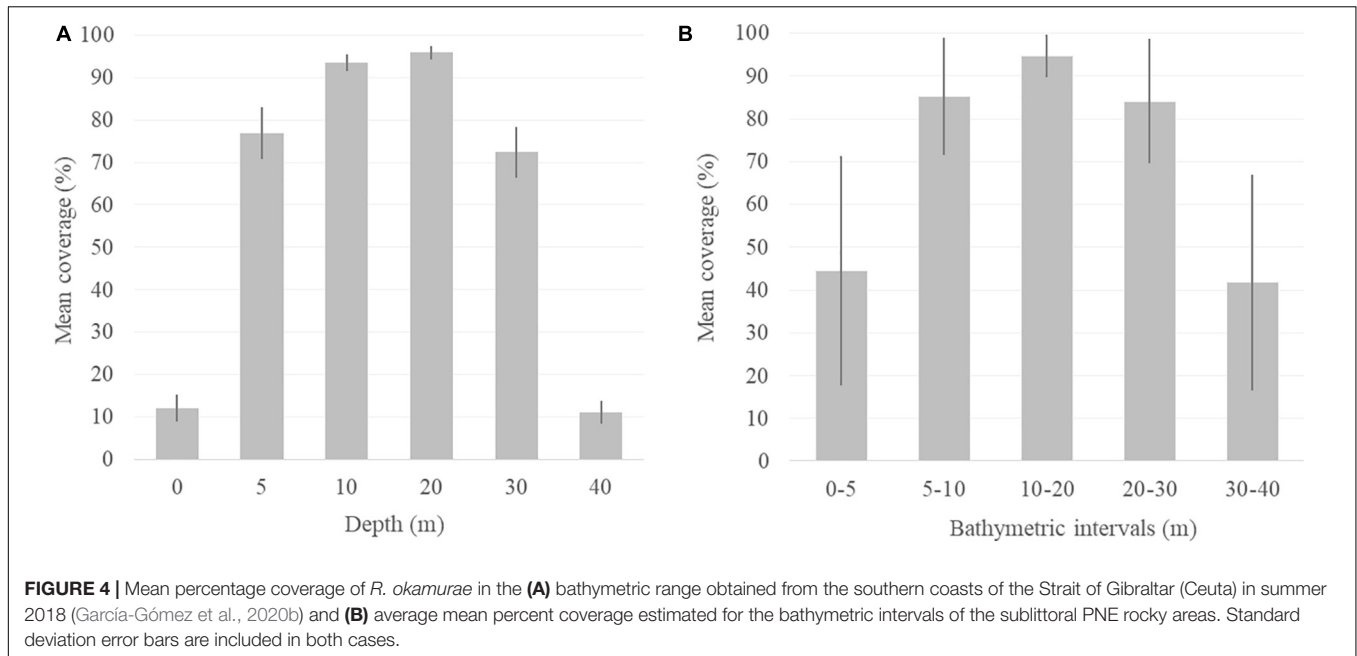
Coverage results obtained in 2018 by García-Gómez et al. (2020b) on horizontal illuminated rocky surfaces, extrapolated here for a larger-scale approximation, suggest a high impact on sublittoral habitats due to the massive occupation of the generalized conquest of rocky areas by *R. okamuræ*. In general, the coverage

data obtained were very high, especially if compared with the biomass data observed in native habitats (Japan) by Agatsuma et al. (2005) (with similar north latitude to Spain, between 20° and 40°). Results obtained for coverage extrapolation coincide with those obtained in 2019–2020 by CAGPyDS (2020) for the bathymetric intervals between 0 and 20 m (80–100% *R. okamuræ* coverage). However, the same results were not found for the intervals between 20 and 30 m (10–50%), so it could be interpreted a possible decline between 2018 and 2020 in the depth ranges where the illumination exposition is lower, so the competitive interactions with the resident macrobiota have not been able to be maintained after the initial stages of strong expansion after 2016.

Following the favorability models (Real et al., 2006), *R. okamuræ* could proliferate within the ecological environments within the Atlantic coasts of Andalusia and Morocco, the Mediterranean Sea and the Black Sea (Muñoz et al., 2019). In fact, it has been confirmed the fast progress of the species toward the Mediterranean and Atlantic waters, and the recent arrival by wrack deposits at the nearby coasts of Granada and Almería (Altamirano et al., 2019; CAGPyDS, 2020; Figueroa et al., 2020). The nature of the ecosystem implications by *R. okamuræ* establishment in the recipient ecosystems have been previously exposed in García-Gómez et al. (2018); Altamirano et al. (2019) and García-Gómez et al. (2020b). Results obtained in these experiences and the spread dynamic of the species alarm the rate of recorded invasions by marine macroalgae that have taken the place of target resident species becoming dominant in the last years (Boudouresque and Verlaque, 2002). This makes the geographic expansion along the rocky surfaces of the PNE littoral worrying and stress the detriment of the benthic biota already attributed to the establishment of *R. okamuræ* in the area. In the southern coasts of the Strait, habitat changes derived from *R. okamuræ* establishment have proved to have implications in endangered coralligenous species (Sempere-Valverde et al., 2020) and associated fauna to resident macroalgae (Navarro-Barranco et al., 2019), so it could be also expected that effects on sessile communities can be also translated to other ecosystem components. As Levine et al. (2004) propose, recognizing that biotic containment can occur through species interactions, it could be expected that ecosystem components interacting with *R. okamuræ* could regulate the invasive populations dominance. In this sense, more efforts are needed studying interspecific interactions involving the invasive species and the incipient role of co-occurring specific resident taxa taking advantage of the spatial colonization [e.g., *A. armata* has been observed on dense populations of *R. okamuræ* (unpublished data)].

Moreover, the high coverage estimations challenge the ecosystem and socio-economic services in the area. The PNE is an integral part of the Mediterranean Intercontinental Biosphere Reserve. It is frequented by tourists and scuba divers, while receives a high influence from both commercial and recreational traffic from areas as Algeciras Bay (Bermejo et al., 2014). Although impacts on socio-economic services have not being investigated, the excess of biomass shortly after *R. okamuræ* establishment reported substantial impacts in the area, both in tourism and fisheries (e.g., the trap of

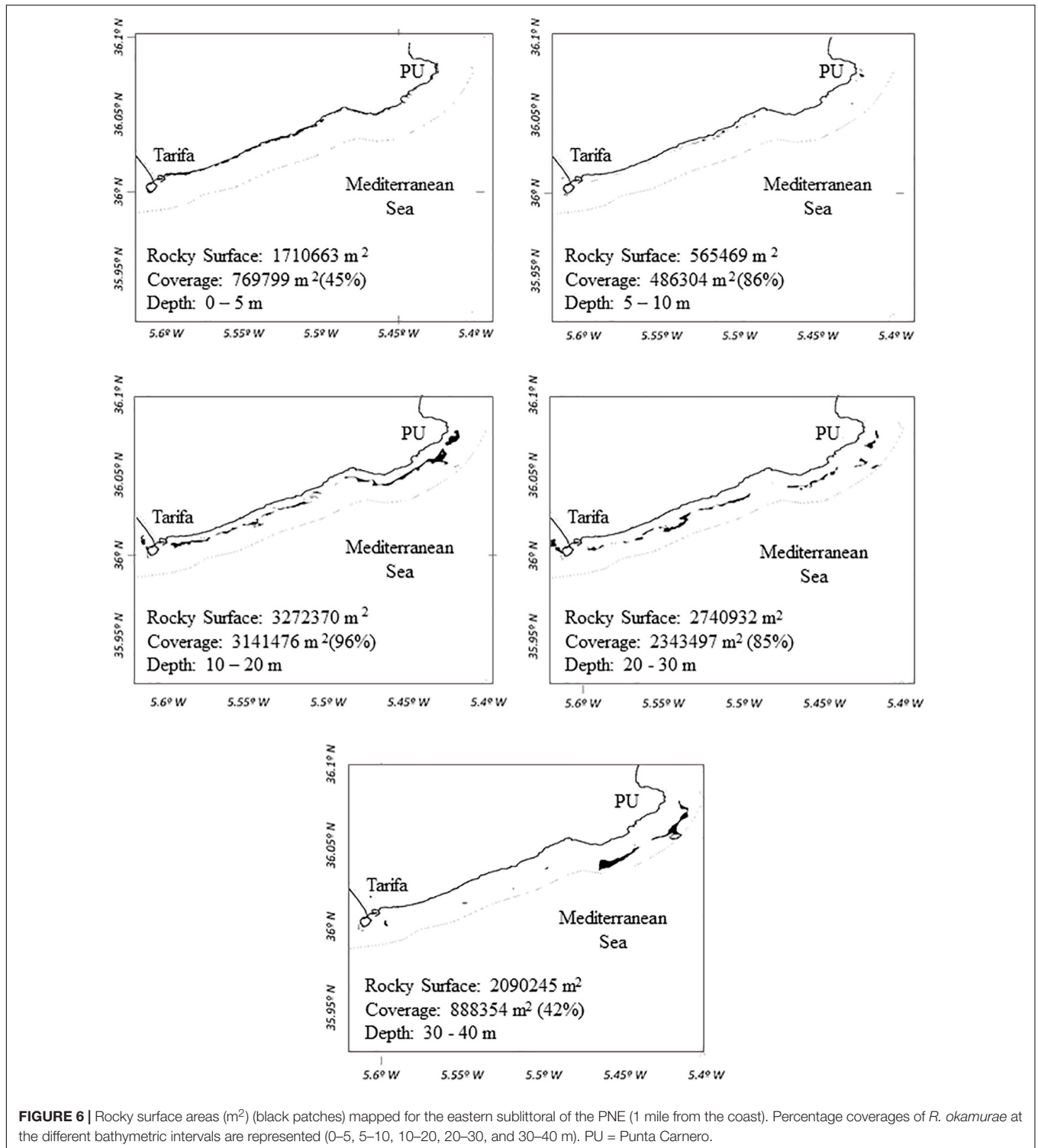




Tarifa) (García-Gómez et al., 2018; Altamirano et al., 2019). If we consider the ability of the species to easily remain attached on nets and other artificial materials (García-Gómez et al., 2018), results obtained for estimations at 1 mile from the coastline enhance the attention on potential impacts in practices developed in the area. Moreover, taking into account that the species was inadvertently introduced via marine aquaculture within the European waters (Thau Lagoon) (Verlaque et al.,

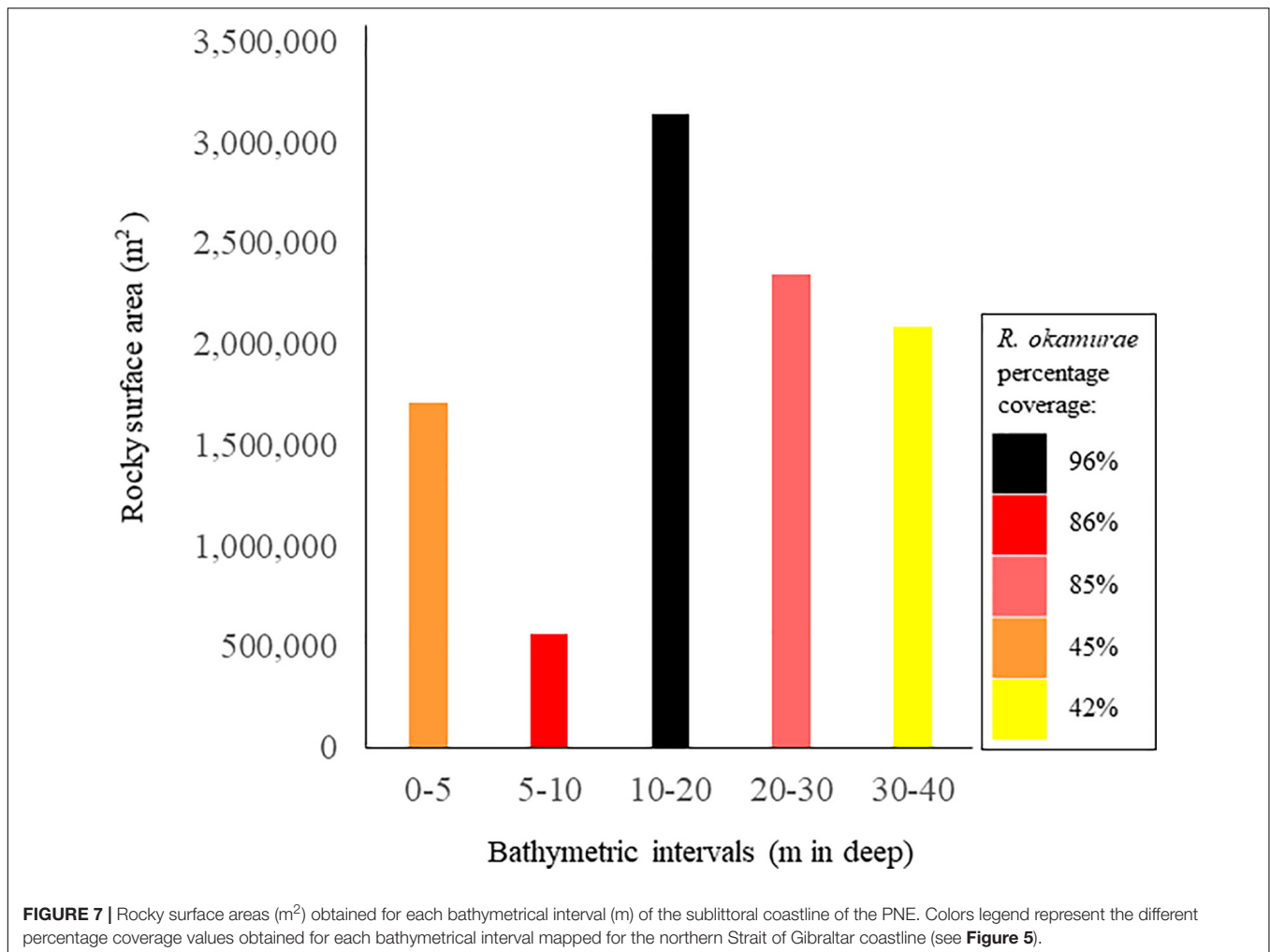
2009), preventive strategies for secondary spread pathways across the Mediterranean and the Atlantic waters must be strongly considered.

Facing with the arduous challenge to strategies applied in advanced invasion stages, mitigation efforts could be essential to protect similar areas not yet impacted but susceptible to be invaded. It is at this point that monitor/modeling techniques play a key role, since estimations based on cartographies can



result useful to identify areas not yet colonized. Moreover, existing precedents of successful actions against invasive marine macroalgae establishment and propagation have revealed that monitoring strategies are not effective unless applied in early stages of colonization, when the species has a limited spatial distribution (Anderson, 2007). In fact, according to Ojaveer

et al. (2015), if the species has already managed to establish in large areas, eradication is unlikely. In this sense, it is worth highlighting the case of *C. taxifolia* as a precedent of marine bioinvasion which invasive process resulted impossible to be stopped by control efforts (Ruesink and Collado-Vides, 2006), becoming the most widespread invasive macroalga in the



Mediterranean waters, occupying 20,000 Ha of sublittoral areas (Anderson, 2007).









Before-After Seascape Scenarios Reflecting Extreme Changes Due *Rugulopteryx okamuræ* Invasiveness

The comparative analyses of images can make it possible to obtain inferences about the behavior of the biota and the ecological connectivity, since the underwater seascape can have a great influence (Grober-Dunsmore et al., 2009). In this regard, the Citizen Science initiative developed in this study allowed a large spatial and temporal dataset to visualize impacts related with *R. okamuræ* establishment, providing an accurate underwater seascape of the bioinvasion consequences. The seascape sampling provided an adequate approach for monitoring developments and it was useful for describing and categorizing some benthic communities that interact with the brown macroalga in the PNE littoral. Comparisons before and after *R. okamuræ* establishment revealed an overall substitution of the benthic seascapes by the invasive species, but also effectively evidenced negative impacts on particular resident species. For example, the disappearance

of sea urchin species [*Arbacia lixula* (Linnaeus, 1758) and *Paracentrotus lividus* (Lamarck, 1816)] from different shallow rocky bottoms could be inferred by comparing pairs of images examined. Indeed, in most of the cases, signs of total colonization were observed only 3 years after its establishment in the Strait of Gibraltar and thus, impacts by the generalized substitution of the resident macroalgae at illuminated and shaded habitats (i.e., native and invasive species already established in the area, as those from the genus *Asparagopsis*) were particularly visible.


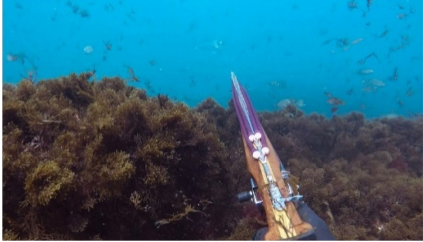





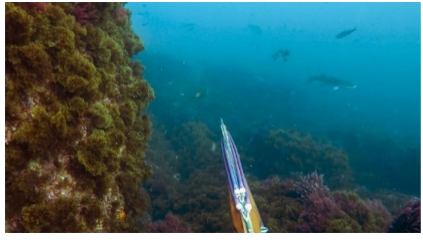
In situ observational data also increased the quantity of *R. okamuræ* observations available for ecological researches (as it has previously been pointed out by Crall et al., 2015). Image data from 2016 at Tarifa Island revealed that *R. okamuræ* monopolized more than 80% of highly illuminated horizontal surfaces at hard bottoms between 5 and 10 m depth, which contrast with results obtained at partially shaded vertical substrata sampled in the SBPQ station in the same year, where <10% mean coverage was estimated by photoquadrat analyses. In the latter habitats, *R. okamuræ* coverage increased later, in 2017 (<60% coverage) (García-Gómez et al., 2020b), and thus revealing habitat-dependent patterns not previously perceived.

TABLE 2 | Comparisons before and after *R. okamurae* first apparition in the Strait of Gibraltar coasts (2015) in locations surveyed in the study area.

Before 2015	After 2015
 <p data-bbox="113 530 766 611">May 2011 Bare horizontal and vertical metallic surfaces (artificial substrata) with associated coverage of photophilic crustose macroalgae</p>	<p data-bbox="715 259 877 279">Caldera Profunda</p>  <p data-bbox="831 530 1476 611">September 2018 Horizontal and vertical metallic surfaces highly colonized by <i>R. okamurae</i></p>
 <p data-bbox="113 924 766 1025">September 2006 Bare horizontal and vertical metallic surfaces constituting the artificial substrata with associated populations of <i>Treptacantha usneoides</i> (Linnaeus) Orellana & Sansón (arborescent talli of large size)</p>	<p data-bbox="715 652 877 673">Tanque Gasoil</p>  <p data-bbox="831 924 1476 1025">September 2016 Horizontal and vertical metallic surfaces constituting the artificial substrata highly coated by <i>R. okamurae</i>. Populations of <i>T. usneoides</i> were not observed</p>
 <p data-bbox="113 1338 766 1440">September 2006 Horizontal illuminated natural habitats colonized by <i>T. usneoides</i> (arborescent thalli), <i>Halopteris scoparia</i> (Linnaeus) Sauvageau, <i>Colpomenia sinuosa</i> (Mertens ex Roth) Derbès & Solier and crustose macroalgae of the genus <i>Lithophyllum</i></p>	<p data-bbox="715 1067 877 1087">La Garita</p>  <p data-bbox="831 1338 1476 1440">September 2016 Horizontal illuminated natural habitats monopolized by <i>R. okamurae</i></p>
 <p data-bbox="113 1742 766 1827">February 2006 Intertidal pool walls subjected to long shadow periods along the day, with no signals of <i>R. okamurae</i> presence</p>	<p data-bbox="715 1481 877 1502">Pozas Intermareales</p>  <p data-bbox="831 1742 1476 1827">May 2019 Overall presence of <i>R. okamurae</i> (discontinuous circle) closed to the vertical walls of the intertidal pools</p>






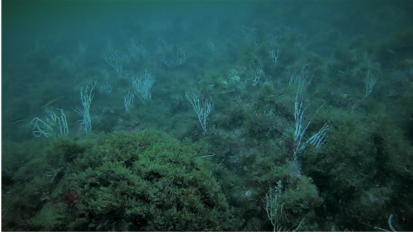
(Continued)

TABLE 2 | Continued

Before 2015	After 2015
 <p data-bbox="113 540 703 617">October 2012. Natural rocky substrata scarcely colonized by resident macroalgae while significant coverages of photophilic crustose macroalgae</p>	<p data-bbox="762 256 831 277">El Tejar</p>  <p data-bbox="831 540 1437 617">October 2018 Natural rocky substrata totally coated by <i>R. okamurae</i> and detached mats floating on the water column</p>
 <p data-bbox="113 934 703 1011">July 2012 The seascape was characterized by the high presence of the invasive resident species <i>A. armata</i> and the sympatric native macroalga <i>H. scoparia</i></p>	<p data-bbox="762 627 839 648">La Caleta</p>  <p data-bbox="831 934 1437 1011">July 2019 <i>R. okamurae</i> monopolized the rocky substrata. Affections by <i>R. okamurae</i> were observed on the photophilic crustose coralline communities</p>
 <p data-bbox="113 1321 703 1369">August 2012 Natural rocky substrata where the sea urchin <i>A. lixula</i> is presented</p>	<p data-bbox="762 1021 836 1042">La Isleta</p>  <p data-bbox="831 1321 1437 1369">August 2018 Surfaces totally coated by <i>R. okamurae</i>, while <i>A. lixula</i> was not detected</p>
 <p data-bbox="113 1680 703 1757">August 2012 Wide coverage of resident macroalgae (mainly native species) on natural rocky substrata</p>	<p data-bbox="762 1415 871 1435">Barranco Hondo</p>  <p data-bbox="831 1680 1437 1813">September 2018 Replacement scenarios of the benthic macrobenthos by <i>R. okamurae</i>, together with the presence of the previously established invasive macroalga <i>A. armata</i>. Affections by <i>R. okamurae</i> were also observed on the photophilic crustose coralline communities</p>

(Continued)

TABLE 2 | Continued

Before 2015	After 2015
<p data-bbox="231 285 646 520"></p> <p data-bbox="113 530 715 629">May 2012 Natural rocky substrata with hardly any coverage of macroalgae and a generalized presence of the sea urchins <i>A. lixula</i> and, to a lesser extent, <i>P. lividus</i></p>	<p data-bbox="735 256 858 273">Guadalmesí I</p> <p data-bbox="948 285 1362 520"></p> <p data-bbox="831 530 1476 607">May 2019 Apparent dominance of <i>R. okamurae</i> and disappearance of a large part of populations of the species <i>A. lixula</i> and <i>P. lividus</i></p>
<p data-bbox="231 679 646 913"></p> <p data-bbox="113 924 715 1029">August 2015 Apparent high abundances of resident macroalgal species (e.g. <i>Dyctiota dichotoma</i> var. <i>intricata</i> (C. Agardh) Greville, <i>Dyctiota fasciola</i> (Roth) J. V. Lamouroux and <i>A. armata</i>)</p>	<p data-bbox="735 650 858 667">Guadalmesí II</p> <p data-bbox="948 679 1362 913"></p> <p data-bbox="831 924 1476 1029">June 2017 Generalized presence of <i>R. okamurae</i> to the detriment of the previously established resident benthic biota. Affections by <i>R. okamurae</i> were also observed on photophilic crustose coralline communities</p>
<p data-bbox="231 1104 646 1338"></p> <p data-bbox="113 1348 715 1425">October 2016 The gorgonian colonies of <i>Eunicella singularis</i> (Esper, 1791) with resident macroalgal associated communities on rocky substrata (20–25 m depth)</p>	<p data-bbox="735 1075 858 1091">Punta Carnero</p> <p data-bbox="948 1104 1362 1338"></p> <p data-bbox="831 1348 1476 1475">September 2019 The macroalgal community was replaced by the monopolized presence of <i>R. okamurae</i>. Adjunct substrata were practically colonized by the invasive macroalga (20–25 m depth) and the spatial pressure on gorgonians present was apparent</p>

CONCLUSION

The present study determined that impacts derived from *R. okamurae* establishment remain high in the rocky habitats studied of the PNE. Monitoring studies on the SBPQ station of Tarifa Island revealed a high spatial establishment of the invasive species since its first detection in 2015 and an efficient loss in the sessile resident biota in the latter years, even in periods of minimum growth. In view of the ecosystem implications, coverage values estimated for rocky habitats (over 85% between 10 and 30 m depth) claim monitoring efforts focused in

threatened habitats not yet colonized a remain step ahead of the drastic scenarios observed. In this regard, Citizen Science collaborations for the detection, evaluation and monitoring of impacts from *R. okamurae* resulted a useful and promising tool for further studies. We consider that these monitoring initiatives would be even more successful if combined with periodic monitoring methodologies under specific designs. Thus, monitoring stations located in areas coinciding with those where contributors act (e.g., SBPQ station in Tarifa Island), could allow citizen collaboration through the applicability of non-invasive tools for image analysis procedures easy to understand and apply.

Only if these tools are promoted within local networks (i.e., high anthropic pressure areas), they could help to early detect and monitor (e.g., “before-after” approximations) local (e.g., urban discharges, oil slicks) or global environmental impacts (e.g., global change), which would facilitate to act in time in the face of bioinvasion schemes.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/**Supplementary Material**, further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS

JG-G involved in investigation, writing – original draft, analysis data, supervision, validation, funding acquisition, and project administration. MF involved in investigation, writing – original draft, analysis data, and data curation. LO-P involved in investigation, special collaboration writing – original draft, and data curation. JRDdR involved in investigation and collaboration in cartography of coverage. ID-A involved in investigation, analysis data and special collaboration writing – original draft. MC, JQ, and SM collaboration underwater images “before-after” impact (Citizen Science). CM involved in investigation and supervision; JG-G, MF, LO-P, ID-A, and CM contributed substantially to revisions of manuscript. All authors contributed to the article and approved the submitted version.

FUNDING

This study was funded by the CEPESA Foundation and Red Eléctrica de España (REE). Additional financial support was provided by the Organization of Fish Producers of Almadrabas (OPP-51), ENDESA, ACERINOX, and Diputación Provincial de Cádiz. This project has also benefited from scientific equipment and infrastructures financed by the AP of Seville and the

Aquarium of Seville and by the VI Plan Propio Universidad de Sevilla via a post-doctoral grant (ID-A).

ACKNOWLEDGMENTS

We are deeply grateful to Manuel Martínez Chacón, Juanjo Quintero, and Salvador Magariño for their selfless collaboration “Citizen Science” in the search, obtaining underwater images, and selection of graphic information, especially from documentary archives prior to 2015. We are also grateful to Red Eléctrica de España (REE) and CEPESA Foundation for mean financial support to scientific research related to the environmental impact of the invasive species *Rugulopteryx okamurae* on the marine autochthonous biodiversity of the Bay of Algeciras, Strait of Gibraltar, and nearby areas, and to Organization of Fish Producers of Almadrabas (OPP-51), ENDESA, ACERINOX, and Diputación Provincial de Cádiz for additional financial support, related with the presence of invasive algae in their artisanal fishing or industrial port infrastructures. We thank the Port Authority of Seville (APS) and Seville Aquarium for scientific infrastructure acquired in project number 1889 (FIUS), related to the estuary of the Guadalquivir River implied in environmental monitoring campaigns and controls in the areas near the mouth of the mentioned river. We thank ESGEMAR, S.A., for providing cartographic support and map delimitation and CIES (Algeciras) and Campo de Gibraltar (La Línea) diving clubs. We also thank Isabel Cárdenas, Curro Sánchez, Rocío Espada, Juan Miguel Recio, Antonio Vegara, José Fosela, and Candela Sánchez for their continued collaboration. This paper has been benefited in the city of La Línea by the logistical support of Puerto La Alcaidesa, the Yacht Club (RCNLL), and the Maritime Club, as well as by the town halls of this city and Tarifa.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fevo.2021.639161/full#supplementary-material>

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Conflict of Interest: JRDdR was employed by the company Esgemar S.A.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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