Article

Impacts of the Invasive Alien Carpobrotus spp. on Coastal Habitats on a Mediterranean Island (Giglio Island, Central Italy)

Michele Mugnai 1,*, Renato Benesperi 1, Daniele Viciani 1,*, Giulio Ferretti 2, Michele Giunti 3, Francesca Giannini 4 and Lorenzo Lazzaro 1

1 Department of Biology, University of Florence, Via La Pira 4, 50121 Firenze, Italy
2 University Museum System, Botanical Garden “Giardino dei Semplici”, University of Florence, Via Micheli 3, 50121 Firenze, Italy
3 Nature and Environment Management Operators s.r.l., Piazza M. D’Azeglio 11, 50121 Firenze, Italy
4 National Park of the Tuscan Archipelago, 57037 Portoferraio, Italy
* Correspondence: michele.mugnai@unifi.it (M.M.); daniele.viciani@unifi.it (D.V.)

Abstract: Carpobrotus acinaciformis and C. edulis are well-known invasive alien plants native to South Africa, whose detrimental effects on native communities are widely documented in the Mediterranean basin and thus largely managed in coastal ecosystems. Most of the literature on these species focuses on their impacts on habitats of sandy coastal dunes, while the effects of Carpobrotus spp. invasion on other habitats such as rocky cliffs and coastal scrubs and garrigues are almost neglected. We present a study case conducted on a small Mediterranean island where Carpobrotus spp. invaded three different natural habitats listed within the Habitat Directive 92/43/CEE (Natura 2000 codes 1240, 1430, and 5320). We surveyed the presence and abundance of native species and Carpobrotus spp. on 44 permanent square plots of 4 m² in invaded and uninvaded areas in each of the three habitats. We found impacts on plant alpha diversity (intended as the species diversity within each sampled plot) in all the habitats investigated in terms of a decrease in species richness, Shannon index, and abundance. Invaded communities also showed a severe change in species composition with a strong homogenization of the floras of the three habitats. Finally, the negative effect of invasion emerged even through the analyses of beta diversity (expressing the species diversity among sampled plots of the same habitat type), with Carpobrotus spp. replacing a large set of native species.

Keywords: biodiversity; community ecology; conservation; endemic species; indicator species; maquis; nestedness; shrubland; turnover; vegetation

1. Introduction

Biological invasions represent one of the most dramatic threats to biodiversity, contributing substantially to the widespread and accelerated decline in Earth’s biodiversity and associated benefits to people from nature [1,2], a situation even more dramatic considering that an increase in the rate of new introductions is foreseen in the future [3]. Invasive alien plants (IAPs) can exert several deleterious impacts on native communities, leading to a local decrease in plant and animal species richness and diversity [4–6]. Indeed, IAPs can induce cascade effects linked to dramatic changes in the structure and function of invaded ecosystems [5,7,8] and resulting in the reduction of the distinctiveness of local biological communities [9]. Islands and coastal mainland regions have been recently individuated as hotspots of established alien species richness across multiple taxonomic groups [10]. In this context, the deleterious effect can be even higher, and IAPs presence and impacts are well documented on Mediterranean island ecosystems [11–15]. Indeed, the Mediterranean basin, with its complex system of archipelagos, islands and islets, represents an important area of species diversity. It is usually reported that Mediterranean basin vascular flora
amount at about 24,000–25,000 species, accounting for 10% of world plant richness, with at least 13,000 endemics, with rates of endemism often exceeding 10%, and sometimes 20%, of local flora (see [12]).

In Europe, IAPs are considered one of the major drivers of changes in natural and semi-natural habitats and their presence increases the probability of unfavorable conservation status of natural habitats [16], being linked to a general deterioration of biodiversity and to the alteration of habitat structure and functions in plant communities [15,17]. Indeed, there is a general awareness of the potential impacts of IAPs on native plant communities and on the habitats of Community Interest listed in the Habitats Directive (Council Directive no. 92/43/EEC, hereafter N2000 habitats), even if for several habitats and species there is lack of direct evidence [18]. The importance of the N2000 network in tackling the risks posed by biological invasions was underlined by the European Commission in the EU 2020 Biodiversity Strategy, and further emphasized in the recent EU 2030 Biodiversity Strategy. However, the N2000 network efficacy in decreasing the vulnerability to invasive alien species is largely still unknown [18,19]. Accordingly, data on the presence and impact of IAS on the N2000 habitats and on the N2000 network are crucial to counter their detrimental impacts [18].

The species of the genus *Carpobrotus* N.E.Br. (Aizoaceae)—in particular *C. acinaciformis* (L.) L.Bolus and *C. edulis* (L.) N.E.Br. and their hybrids—are considered among the most abundant and most investigated IAPs in the whole Mediterranean basin, where they display a very high invasive potential [12,15,20]. The invasion by *Carpobrotus* spp. causes strong negative impacts on the ecology of invaded ecosystems, mainly sand dunes and rocky sea cliffs [20], with significant changes in the invaded ecosystems at a variety of scales [21–24]. Impacts have been well depicted on several ecosystem components and processes, from plant biodiversity and vegetation structure [13,21,25–28] to soil conditions and physico-chemical and biological processes [28–31], these latter resulting also in the reduction in seed germination and survival of seedlings of the native plants [30,32,33]. In the whole Mediterranean basin, these species have been targeted by several projects of control, mostly fostered by local stakeholders, but also often linked to the EU LIFE program (see also [20], and especially on coastal dunes in the Mediterranean basin *Carpobrotus* is the genus with the largest number of records of control actions [12,34]. In Italy *Carpobrotus* spp. are considered invasive and among the most threatening invasive plants [15], with well-documented impacts at the community level and on N2000 habitats, but with evidence mostly restricted to sand dune habitats (particularly N2000 habitats 2120 “Shifting dunes along the shoreline with *Ammophila arenaria* (white dunes)”, 2210 “*Crucianellion maritimae* fixed beach dunes” and 2250* “Coastal dunes with *Juniperus* spp.”). However, these are not the only habitat types invaded by these IAPs in Italy: in Tuscany, it is particularly invasive in the Tuscan Archipelago [35], where it is one of the most harmful invasive alien species, particularly affecting the rocky cliffs coastal vegetation of the islands of the Archipelago [14].

Hence, studies on the impacts related to IAPs are very important as they represent a valuable source of information necessary to lay the basis for any generalization on the scenario of biological invasions and are an important tool to implement and enforce more effective management strategies [2,36]. To provide a more comprehensive understanding of impacts, composite approaches are needed, assessing the effects of invasive plants on several response variables [36]. Moreover, to provide a better comprehension of the ecological process caused by IAPs invasion, the study of beta diversity could provide important insights. Beta diversity is defined as the ratio between gamma (regional) and alpha (local) diversities, and it essentially quantifies the number of different communities in the region [37]. Indeed, beta diversity studies can provide considerable insights into the importance of deterministic and stochastic processes in generating community structure along spatial and ecological gradients [37,38], and have been already used to provide insights into drivers and mechanisms of invasion and assembly of alien communities at a broad spatial scale [39,40]. Decomposing beta diversity in different components (i.e.,
Plants 2022, 11, 2802

...disentangling the contribution of spatial turnover and nestedness to beta diversity patterns) allows us to highlight patterns linked to the biological phenomena in the act and is essential for analyzing the causality of the processes underlying biodiversity [37].

Within this work, we aimed to assess the impact of *Carpobrotus* spp. on the native communities at Giglio Island (Tuscan Archipelago, Italy), providing a baseline of data on the impacts of these IAPs on invaded ecosystems as pre-intervention monitoring linked to the control of *Carpobrotus* spp. foreseen within the LIFE project LETSGO GIGLIO “Less alien species in the Tuscan Archipelago: new actions to protect Giglio island habitats” ([www.lifegogiglio.eu](http://www.lifegogiglio.eu), accessed on 16 October 2022). Moreover, we aimed at providing comprehension of ecological processes in the act to better inform conservation and restoration efforts. Accordingly, we aimed to (i) verify the impact of *Carpobrotus* spp. on native species richness, diversity, and total cover, (ii) evaluate the impacts on the native species composition of invaded communities, and (iii) assess the main processes in the act (turnover vs. species loss) focusing on beta diversity features of invaded and non-invaded communities. Toward these aims, we monitored a series of vegetation plots within the three main coastal habitats invaded by *Carpobrotus* spp. on Giglio island (N2000 habitats 1240: vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp., 1430: Halo-nitrophilous scrubs (*Pegano-Salsoletea*) and 5320: Low formations of *Euphorbia* close to cliffs).

2. Results

The sampling led to the identification of 74 species in 88 plots (see Supplementary Tables S1 and S2). Vegetating cover varied substantially in Habitat 1240 between Control and Invaded plots, while it was more consistent in Habitats 1430 and 5320, even if in these habitats there was a high variation between 2020 and 2021 (see Supplementary Table S3). The results generally showed in all three habitats the presence of significant impacts on alpha diversity linked to *Carpobrotus* spp. invasion (Table 1).

*Carpobrotus* spp. cover was stable in 2020 and 2021 around 60% in habitat 1240 and varied significantly from about 50% in 2020 to 25% in 2021 in habitat 1430 and from about 75% in 2020 to 50% in 2021 in habitat 5320 (Figure 1A). Noteworthy, a few *Carpobrotus* spp. seedlings arrived in 2021 in four control plots: two in habitat 1430 and two in habitat 5320 (see zoomed bars in Figure 1A). The *Carpobrotus* spp. dead litter showed an inverse variation, being more or less stable around 20% in habitat 1240 and increasing in 2021 in both habitat 1430 and habitat 5320 (Figure 1B).

Table 1. Repeated Measurement ANOVA table for the effect of Invasion Status (Control plots vs. Invaded plots) and Year (sampling year 2020 vs. 2021) on Native species total cover, native species richness and native species diversity expressed as H’, provided for each Natura 2000 habitat. N2000 habitat codes: 1240 = vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp., 1430 = Halo-nitrophilous scrubs (*Pegano-Salsoletea*) and 5320 = Low formations of *Euphorbia* close to cliffs. numDF: numerator degree of freedom; denDF: denominator degree of freedom. Significance codes: *p value < 0.001 ‘***’; *p value < 0.01 ‘**’; *p value < 0.05 ‘*’; *p value < 0.10 ‘.’.

| Response Variable | Habitat | Variable         | numDF | denDF | F Value | p Value |
|-------------------|---------|------------------|-------|-------|---------|---------|
| Native species total cover | 1240 | Invasion Status | 1     | 44    | 155.35  | <0.001 *** |
|                   |        | Year             | 1     | 44    | 0.25    | 0.616 |
|                   |        | Invasion Status:Year | 1     | 44    | 0.54    | 0.466 |
|                   | 1430 | Invasion Status | 1     | 12    | 19.69   | <0.001 *** |
|                   |        | Year             | 1     | 12    | 4.46    | 0.056 |
|                   |        | Invasion Status:Year | 1     | 12    | 0.15    | 0.705 |
|                   | 5320 | Invasion Status | 1     | 20    | 101.44  | <0.001 *** |
|                   |        | Year             | 1     | 20    | 0.13    | 0.720 |
|                   |        | Invasion Status:Year | 1     | 20    | 9.21    | 0.007 *** |

* Variables log transformed.
Table 1. Cont.

| Response Variable | Habitat | Variable | numDF | denDF | F Value | p Value |
|-------------------|---------|----------|--------|--------|---------|---------|
|                   | 1240 $^£$ | Invasion Status | 1 | 44 | 8.32 | 0.006 ** |
|                   |         | Year | 1 | 44 | 0.45 | 0.504 |
|                   |         | Invasion Status:Year | 1 | 44 | 0.11 | 0.743 |
| Native species richness | 1430 | Invasion Status | 1 | 12 | 18.24 | 0.001 ** |
|                   |         | Year | 1 | 12 | 0.51 | 0.487 |
|                   |         | Invasion Status:Year | 1 | 12 | 0.80 | 0.388 |
|                   | 5320 $^£$ | Invasion Status | 1 | 20 | 35.09 | <0.001 *** |
|                   |         | Year | 1 | 20 | 15.75 | <0.001 *** |
|                   |         | Invasion Status:Year | 1 | 20 | 8.12 | 0.010 * |
| Species diversity (H') | 1240 $^£$ | Invasion Status | 1 | 44 | 3.93 | 0.054 ˚ |
|                   |         | Year | 1 | 44 | 4.80 | 0.034 * |
|                   |         | Invasion Status:Year | 1 | 44 | 0.29 | 0.595 |
|                   | 1430 | Invasion Status | 1 | 12 | 3.46 | 0.088 ˚ |
|                   |         | Year | 1 | 12 | 0.15 | 0.703 |
|                   |         | Invasion Status:Year | 1 | 12 | 9.29 | 0.010 * |
|                   | 5320 | Invasion Status | 1 | 20 | 93.61 | <0.001 *** |
|                   |         | Year | 1 | 20 | 3.23 | 0.087 ˚ |
|                   |         | Invasion Status:Year | 1 | 20 | 0.74 | 0.400 |

$^£$ Variables log transformed.

Figure 1. Cover of (A) *Carpobrotus* spp. and (B) *Carpobrotus* spp. dead litter in the sampled plot according to N2000 habitat, status of invasion (C = Control plots, I = Invaded plots) and year. N2000 habitat codes: 1240 = vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp.,
1430 = Halo-nitrophilous scrubs (*Pegano-Salsoletea*) and 5320 = Low formations of *Euphorbia* close to cliffs. Error bars correspond to standard error. Asterisks indicate significant differences between control and invaded plots, [when in square bracket indicate significant differences within the interaction term Invasion Status:Year]. Significance codes: \( p \) value < 0.001 ‘***’; \( p \) value < 0.01 ‘**’; \( p \) value < 0.05 ‘*’. In lower panel of (A), the *Carpobrotus* cover scale is highly magnified to allow the reading of very small values of cover.

We detected highly significant and marked loss of total abundance of native species in the plots monitored in all three habitats (Figure 2A, Table 1). As to the impacts on alpha diversity, we detected a significant decrease in native plant species richness in the three habitats (Figure 2B, Table 1), with habitat 1240 and habitat 1430 showing an effect irrespective of the year of sampling, while for habitat 5320, the effect varied slightly in the two survey years, with a smaller difference in 2021.

Figure 2. (A) Native species richness, (B) native species diversity expressed as Shannon index \( H' \) and (C) native species abundance (expressed as sum of percentage cover of each species) in the sampled plot according to N2000 habitat, status of invasion (C = Control plots, I = Invaded plots) and year. N2000 habitat codes: 1240 = vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp., 1430 = Halo-nitrophilous scrubs (*Pegano-Salsoletea*) and 5320 = Low formations of *Euphorbia* close to cliffs. Error bars correspond to standard error. Asterisks indicate significant differences between control and invaded plots, [when in square bracket indicate significant differences within the interaction term Invasion Status:Year]. Significance codes: \( p \) value < 0.001 ‘***’; \( p \) value < 0.01 ‘**’; \( p \) value < 0.05 ‘*’, \( p \) value < 0.10 ‘˙’.
With regard to Shannon (H') diversity (Figure 2C, Table 1), while habitat 1240 did not show any significant difference among invaded and control plots, habitat 5320 showed a significant decrease in diversity in the invaded areas irrespective of the year of sampling. For habitat 1430, on the other hand, the effect varies in the two survey years, with a comparable diversity in 2021 between controls and invaded areas.

From the compositional point of view, both the ISA (Table 2) and the NMDS (Figure 3) testify the important changes linked to the invasion by *Carpobrotus* spp.

**Table 2.** Results of the Indicator Species Analysis according to N2000 habitat and invasion status (Control plots vs. Invaded plots). N2000 habitat codes: 1240 = vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp., 1430 = Halo-nitrophilous scrubs (*Pegano-Salsoletea*) and 5320 = Low formations of *Euphorbia* close to cliffs. Significance codes: p value < 0.001 ***; p value < 0.01 **; p value < 0.05 *.

| Species | Invasion Status | Habitat | Indicator Value | p Value |
|---------|----------------|---------|----------------|---------|
| *Limonium planesiae* Pignatti | Control plots | 1240 | 0.95 | <0.001 *** |
| *Brachypodium retusum* (Pers.) P.Beauv. | Invaded plots | 1430 | 0.59 | <0.001 *** |
| *Artemisia arborescens* (Vaill.) L. | 5320 | 0.39 | 0.028 * |
| *Lysimachia arvensis* (L.) U.Manns & Anderb. | 1240 | 0.95 | <0.001 *** |
| *Helichrysum litterum* Guss. | 5320 | 0.39 | 0.028 * |
| *Paronychia echinulata* Chater | 1430 | 0.70 | <0.001 *** |
| *Linum trigynum* L. | 1240 | 0.52 | 0.001 *** |
| *Festuca danthonii* Asch. & Graebn. subsp. *danthonii* | 1430 | 0.50 | <0.001 *** |
| *Lotus hirsutus* L. | 1240 | 0.50 | <0.001 *** |
| *Hypocharis glabra* L. | 1240 | 0.49 | <0.001 *** |
| *Arisorum vulgare* O.Targ.Tozz. subsp. *vulgare* | 1430 | 0.39 | <0.001 *** |
| *Myrtus communis* L. | 1240 | 0.39 | <0.001 *** |
| *Plantago lanceolata* L. | 1240 | 0.39 | <0.001 *** |
| *Stachys major* (L.) Bartolucci & Peruzzi | 1430 | 0.39 | <0.001 *** |
| *Sedum rubens* L. | 1430 | 0.39 | <0.001 *** |
| *Muscaria comosum* (L.) Mill. | 1240 | 0.39 | <0.001 *** |
| *Cytisus laniger* DC. | 1240 | 0.39 | <0.001 *** |
| *Allium roseum* L. subsp. *roseum* | 1240 | 0.39 | <0.001 *** |
| *Euphorbia segetalis* L. | 1240 | 0.39 | <0.001 *** |
| *Lotus edulis* L. | 1240 | 0.39 | <0.001 *** |
| *Polycarpum tetraphyllum* (L.) L. | 1240 | 0.39 | <0.001 *** |
| *Hypocharis acharophorus* L. | 1240 | 0.39 | <0.001 *** |
| *Carlina corymbosa* L. | 1240 | 0.39 | <0.001 *** |
| *Clebionis segetum* (L.) Fourn. | 1240 | 0.39 | <0.001 *** |
| *Urospermum picroides* (L.) Scop. ex F.W.Schmidt | 1240 | 0.39 | <0.001 *** |
| *Runex bucephalophorus* L. | 1240 | 0.39 | <0.001 *** |
| *Coleostephus myconis* (L.) Cass. ex Rchb.f. | 1240 | 0.39 | <0.001 *** |
| *Valantia muralis* L. | 1240 | 0.39 | <0.001 *** |
| *Lolium rigidum* Gaudin | 1240 | 0.39 | <0.001 *** |
| *Jacobaea maritima* (L.) Pelser & Meijden subsp. *maritima* | 1240 | 0.39 | <0.001 *** |
| *Silene gallica* L. | 1240 | 0.39 | <0.001 *** |
| *Echium plantagineum* L. | 1240 | 0.39 | <0.001 *** |
| *Avena barbata* Pott ex Link | 1240 | 0.39 | <0.001 *** |
| *Pistacia lentiscus* L. | 1240 | 0.39 | <0.001 *** |
| *Anisantha madritensis* (L.) Nevski subsp. *madritensis* | 1240 | 0.39 | <0.001 *** |
| *Rostraria cristata* (L.) Tzvelev | 1240 | 0.39 | <0.001 *** |
| *Plantago bellardii* All. | 1240 | 0.39 | <0.001 *** |
| *Calendula arvensis* (Vaill.) L. | 1240 | 0.39 | <0.001 *** |
| *Dactylis glomerata* L. | 1240 | 0.39 | <0.001 *** |
| *Briza maxima* L. | 1240 | 0.39 | <0.001 *** |
| *Cynodon dactylon* (L.) Pers. | 1240 | 0.39 | <0.001 *** |
Anisantha madritensis (L.) Nevski subsp. madritensis 0.27 0.018 * 
Rostraria cristata (L.) Tzvelev 0.25 0.024 * 
Plantago bellardii All. 0.23 0.028 * 
Calendula arvensis (Vaill.) L. 0.17 0.049 * 
Dactylis glomerata L.

Invaded plots 1430 0.59 <0.001 *** 
Briza maxima L. 0.34 0.005 ** 
Cynodon dactylon (L.) Pers. 0.26 0.008 ** 
Carpobrotus spp. 5320 0.39 <0.001 *** 

Figure 3. Non-metric multidimensional scaling (NMDS) ordination plot based on Bray–Curtis dissimilarities of the 88 sampled plots. Plots are grouped according to N2000 habitat and invasion status (Control plots vs. Invaded plots). Ellipses represent the standard deviation of sampled plot positions. N2000 habitat codes: Ha_1240 (circles) = vegetated sea cliffs of the Mediterranean coasts with endemic Limonium spp.; Ha_1430 (triangles) = Halo-nitrophilous scrubs (Pegano-Salsoletea) and Ha_5320 (squares) = Low formations of Euphorbia close to cliffs. See Supplementary Table S4 for plant full names. Invasion Status codes: Con (red)= Control plots; Inv (blue) = Invaded plots.

The ISA showed a dramatic drop in indicator species in the invaded plots compared to the controls, also testifying important compositional differences among the three habitats in their natural conditions (i.e., in the control plots). The habitat 1240 control plots had only one indicator species (Limonium planasie) while the other two habitats showed a very high number of indicator species in the controls (16 and 21, respectively, in habitat 1430 and habitat 5320). In the invaded status, all three habitats showed very few indicator species, with Carpobrotus spp. being the indicator species in only habitat 5320. The NMDS analysis (stress = 0.092, non-metric fit R² = 0.991, linear fit R² = 0.964) highlighted a well-defined differentiation (as expected) between the control plots of the three habitats, which lay on the upper area of the plots. Conversely, the communities appear closer when invaded by Carpobrotus spp. (lower part of the plot), and particularly habitat 5320 and habitat 1240 invaded plots almost overlap, while habitat 1430 invaded plots are closer to the respective control plots.

Through the assessment of beta diversity, we found different patterns for the three habitats considered. Within habitats 1240 and 5320, in both years we found that the higher levels of beta diversity are between invaded and not-invaded communities, with the greater portion due to species turnover (Figure 4).
defined differentiation (as expected) between the control plots of the three habitats, which lay on the upper area of the plots. Conversely, the communities appear closer when invaded by *Carpobrotus* spp. (lower part of the plot), and particularly habitat 5320 and habitat 1240 invaded plots almost overlap, while habitat 1430 invaded plots are closer to the respective control plots.

Through the assessment of beta diversity, we found different patterns for the three habitats considered. Within habitats 1240 and 5320, in both years we found that the higher levels of beta diversity are between invaded and not-invaded communities, with the greater portion due to species turnover (Figure 4).

Regarding beta diversity within habitat 1430, we found more comparable patterns of diversity within invaded, control, and between control and invaded communities: for 2020, the comparison of control−invaded results were slightly higher, while for 2021, diversity within invaded communities it was higher. In those three cases of comparison (for both 2020 and 2021), the main contribution to beta diversity of habitat 1430 is due to species turnover.

3. Discussion

Our results provide significant evidence on the impacts of *Carpobrotus* spp. on the rocky cliff coastal habitats of Giglio island and offer interesting insight into the ecological processes acting on the invaded sites. The very high impact of *Carpobrotus* spp. at the alpha diversity level has been frequently reported, inducing a decrease in species richness [22,24,26,41,42] of the $H'$ diversity index [26,41,42] and of native plants biomass [22]. However, most of the literature highlighting the general trend of native communities’ depletion when invaded by *Carpobrotus* spp. is focused on sandy dune habitats (see [15,20]) while the impacts on
sea cliff communities are still rarely studied compared to the other types of coastal habitats. Our case brings evidence on the impacts exerted on rocky cliff habitats and vegetation (represented by the mosaic of habitats 1240 and 5320) and coastal cliff shrub vegetation (habitat 5320). Our results are in line with the findings of Buisson et al. [43], who found strong differences in species richness and community composition before and after its removal. We found some differences in the magnitude of the impact. Particularly to 1240, the depletion was less though compared to the other habitats considered, probably as a consequence of the pauci-specific communities typical of this habitat [44]. However, the impact registered is still noteworthy since the poor flora of such habitat includes endemic and high conservation value species such as Limonium sommerianum [44]. The stronger impact of Carpobrotus spp. on native communities for habitat 1430 and habitat 5320 is therefore connected to the normal greater specific richness (note that habitat 1430 is represented on Giglio Island by communities of the syntaxonomic alliance Artemision arborescentis) which characterizes them, increasing the effect of the species disappearance caused by Carpobrotus spp. The results obtained for those habitats are therefore in line with the findings of studies addressing the impact on scrub vegetation [43,45]. Again, however, the research effort focused on sand dune habitat, making the results presented with this study as an update on the knowledge on this topic recognizing the negative impact of Carpobrotus spp. on Mediterranean coastal scrubs and garrigues. We note also that the alpha diversity trends registered slightly differ in the two years of sampling. This phenomenon is probably due to the variation in Carpobrotus spp. cover in the two years (lower in 2021). Indeed, we recorded a reduction in Carpobrotus spp. cover due to the desiccation of some fresh branches (with a correlated increase in dead litter) probably as a consequence of the harsher climatic condition (in particular very low precipitation) to which communities investigated were subjected in the summer of 2020 and spring of 2021 (see Supplementary Table S5). In this year, in fact, the particular aridity may have decreased the diversity in the natural control habitats (due to a lack of annual species) and reduced the dominance of Carpobrotus spp. (which was less abundant) in invaded ones. In particular, habitat 1430, being more structured (more presence of scrubby species) and generally suffering a slightly less pronounced impact than habitat 5320, recorded a greater coverage of native species in the invaded areas than in 2020, with a correlated increase in community diversity.

Our results demonstrate that the invasion of Carpobrotus spp. affects negatively rocky coastal habitats not only from the quantitative point of view (i.e., alpha diversity) but also from the compositional one. Carpobrotus spp. dramatically induced a change in the composition of invaded communities, which resulted as shifted (especially communities of habitat 1240 and habitat 5320) from the not-invaded ones, toward smaller sets of species. The presence of many typical species of the sampled habitats as indicator species only in the not-invaded communities indicate their almost complete depletion in the invaded ones of the same habitat typology, testifying the very important level of replacement caused by Carpobrotus spp. invasion. This confirms that its successful establishment probably operates through the replacement and exclusion of native species, rather than coexistence [15,20]. Moreover, as shown in the NMDS analyses, it is worth mentioning that the communities invaded by Carpobrotus spp. are characterized by the presence of some more ruderal and nitrophilous species (e.g., Cynodon dactylon (L.) Pers. and Mercurialis annua L.). This effect, even if in our case not so pronounced, is comparable with the one observed by Buisson et al. [43] for rocky habitats and by other authors for sandy ones [22]. Indeed, it has already been demonstrated that Carpobrotus spp. may favor the replacement of native plants by ruderal nitrophilous species through soil nutrient enrichment [41,46], hence also in our case the presence of ruderal species could be linked to the significant presence of Carpobrotus spp. litter (as shown in Figure 1B). Moreover, despite the three habitats being naturally characterized by completely separated sets of species, Carpobrotus spp. induced a strong homogenization of their flora. In fact, it has been frequently reported that serious invaders may directly and quickly lead to a strong biotic homogenization of plant communities [47–51]. The impact of
Carpobrotus spp. is an example of this phenomenon, as demonstrated by this study and by others on dune habitats [22].

The analysis of beta diversity components of diversity allowed us to confirm the insight from compositional analyses, shedding light on the main ecological processes in action due to the invasion. For habitat 1240, the higher beta diversity between invaded and not-invaded communities is primarily a consequence of the reduced number of species occurring in contexts impacted by Carpobrotus spp., confirming the results of alpha diversity. The depletion of invaded communities, in terms of the number of species, is responsible for the high beta diversity between invaded and not invaded communities being due mainly to the turnover of species, with Carpobrotus spp. replacing (in many cases entirely) and out-competing native species. This evidence confirms the already shown phenomenon of a shift in community composition due to Carpobrotus spp. (e.g., [20]), even considering the beta diversity. This pattern has been found also for habitat 5320, in which the diversity between invaded and not-invaded communities was considerably higher than that occurring within them. Hence the impact of Carpobrotus spp. induces even in garrigue a severe change in plant communities, leading them to be considerably differentiated from the native ones. In this case, we note a significant difference in the contribution of turnover and nestedness to beta diversity for the two years. As already stated for other results, this effect might be attributed to the drier condition of 2021 and to the consequent reduction in Carpobrotus spp. coverage and the slightly higher affirmation of a few native species more resistant to aridity (e.g., Euphorbia seguieriana Neck. and Helichrysum litoreum Guss.). Consequently, in terms of beta diversity, this entails that in 2020, the diversity between invaded and not-invaded communities is almost entirely due to species turnover (Carpobrotus spp. replacing a large set of native species), while in 2021, the diversity is partially due to nestedness (as the few native species in invaded communities consist of subsets of the invaded one). Finally, the comparable patterns of beta diversity between and within invaded and not-invaded communities can be explained by the intrinsic heterogeneity of this habitat. Despite at the alpha level the effect of Carpobrotus spp. invasion is evident through the reduction in diversity, at the beta diversity level it is less readable, since the diversity is relatively high even within invaded and not-invaded patches. Even in this case the harsher condition of 2021 might have influenced the patterns, which in this case led to a slightly higher beta diversity within invaded communities, probably as a consequence of the less abundance of Carpobrotus spp. in favor of a small recolonization by native species. To our knowledge, the impacts of invasive species have been taken into account addressing almost exclusively the alpha component of diversity, with few examples of studies including the beta diversity component [40,52], of which only one addresses Carpobrotus spp. invasion in coastal areas [53].

In conclusion, within this study, we verified the deleterious impacts on native plant communities linked to the invasion of Carpobrotus spp. on the coastal habitats of a small Mediterranean island (N2000 habitats: 1240 = vegetated sea cliffs of the Mediterranean coasts with endemic Limonium spp., 1430 = Halo-nitrophilous shrubs (Peganum-Salsaletea) and 5320 = Low formations of Euphorbia close to cliffs). These impacts spanned from the decrease in species richness, diversity and abundance to a compositional shift in invaded communities, which also emerged through beta diversity, with Carpobrotus spp. replacing a large set of native species. Indeed, as already shown in several other cases, our data showed that the replacement and exclusion of native species typical of the natural, rather than coexistence, are the main ecological processes linked to Carpobrotus spp. invasion. These results allowed us to outline the current impact of Carpobrotus spp. on the plant communities of Giglio Island, constituting an important baseline of data in view of the interventions aiming to control this invasive species foreseen within project LIFE LETSGO GIGLIO. Moreover, we confirmed the impacts of Carpobrotus spp. invasion on habitats less frequently mentioned in the literature, such as sea rocky cliffs such as 1240 and 5320, and also more structured habitats such as 1430.
4. Materials and Methods

4.1. Study Area

This study took place on the Island of Giglio (WGS84: 42.35527° N, 10.90134° E), which, with its 21.2 km², is the second largest island in the Tuscan Archipelago (Tyrrhenian Sea, Italy) and is located about 14 km in front of Monte Argentario promontory in south Tuscany (Figure 5). The island’s territory is predominantly mountainous, with generally very steep slopes and extensive stretches of denuded rock both inland and along the coast. The climate is Mediterranean, with mild, rainy winters and hot, dry summers peaking in July and August, followed by an autumn resumption of rainfall [54].

Figure 5. Area of study and distribution of the 44 sampling plots according to N2000 habitat and status of invasion. N2000 habitat codes: 1240 = vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp., 1430 = Halo-nitrophilous scrubs (*Pegano-Salsoletea*) and 5320 = Low formations of *Euphorbia* close to cliffs.

The vegetation is purely Mediterranean with the presence of more or less recent holm oak woods and various types of thickets and garrigue [54]. The abandonment of traditional agricultural practices and the shift towards a tourist-based economy has led to a major transformation of the vegetation landscape over the last 70 years, with an increase in the number of thickets and scrub encroaching on former crops (although the last decade has seen an increase in the area returned to cultivation), and an increased impact on coastal habitats [44,54].

Giglio Island is almost entirely included in the Isola del Giglio SAC/SPA (IT51A0023), which covers about 21 km², while the area included within the boundaries of the Tuscan Archipelago National Park is smaller (8.9 km²). The only three small towns remain outside the SAC/SPA: Giglio Castello, Giglio Porto, and the hamlet of Campese, home to the approximately 1550 inhabitants living on the island.
The rocky cliffs coastal vegetation of the island is of particular interest for this study, being the one invaded by *Carpobrotus* spp. and hosts a mosaic of habitats of conservation interest according to Directive 92/43/EEC “Habitat” including the habitat of vegetated sea cliffs of the Mediterranean coasts with endemic *Limonium* spp. (habitat code 1240 according to Dir. 92/43/EEC, and including the important endemic *Limonium sommerianum* Fiori, see also [55], the habitat of Halo-nitrophilous scrubs (*Peganolo-Salsoletea*) (habitat code 1430, represented at Giglio Island by the subtypes identified with the alliance *Artemision arborescentis*) and the habitat of Low formations of *Euphorbia* close to cliffs (hab. code 5320). Particularly, monitoring has been carried out in coastal vegetation at two specific sites strongly invaded by *Carpobrotus* spp.: the promontory of “Punta Capel Rosso” (the island’s southern tip, where *Carpobrotus* spp. is invading two main types of habitats: 1240 and 5320) and the promontory of “Punta del Fenaio” (the island’s northern tip, where *Carpobrotus* spp. is invading two main types of habitats: 1240 and 1430).

4.2. Data Collection

The survey of the vegetation was performed at two locations: Capel Rosso and Fenaio, and was stratified according to the EU habitat, as mapped according to the HaSCITu (Habitat in the Sites of Conservation Interest in Tuscany) program (http://www.regione.toscana.it/-/la-carta-degli-habitat-nei-siti-natura-2000-toscani, accessed on 16 October 2022). During the first phases of the project (between February and late May 2020), several inspections of the islands allowed very detailed mapping of the distribution of *Carpobrotus* spp. in the study area. The experimental monitoring design involves the floristic survey of 2 × 2 m square plots in invaded and control areas. The plots are permanent and were positioned according to a stratified random design based on the surface area of the habitats affected by the actions. For each invaded plot, a paired control as close as possible was selected. Specifically, 44 permanent plots were placed (12 × 2 plots for Habitat 1240, 6 × 2 plots for Habitat 5320, and 4 × 2 plots for Habitat 1430). The higher number of replies for habitat 1240 is linked to the higher surface occupied by this habitat (and invaded by *Carpobrotus* spp.) in the area of study. Each plot was georeferenced, and vegetation sampling was carried out during the vegetative period (May–June) in 2020 and 2021: these two years represent the baseline of data for long-term monitoring of the future interventions of removal and will be carried out for the following years after the interventions to observe the evolution of the vegetation. In each plot, we collected information on the cover of the fresh *Carpobrotus* spp. mat and of its dead litter, as well as of each native species using a percentage scale, taking into account the overlapping of different species (total cover was recorded). Repeated sampling over 2 years led to the survey of 88 plots.

4.3. Statistical Analyses

The effects of the invasion by *Carpobrotus* spp. on alpha diversity of native vegetation, and particularly on the habitats worthy of conservation 1240, 1430, and 5320 were evaluated using Repeated Measurement ANOVA-type models fitting a series of Generalized Least Squares models (GLSm), accounting for a Gaussian spatial correlation of the observations (linked to both the paired structure of sampling design and the presence of two separated localities) and taking also into account that the same plot was revisited for the two years of sampling. For each of the three habitats separately, we assessed whether native species richness (SR), native species diversity expressed as *H*’ index and native species abundance (expressed as the sum of percentage cover of each species) varied according to the status of invasion (invaded vs. control plots). The response variables were log-transformed, when needed, to achieve normality of residuals.

We studied the changes in the species composition of plots using multivariate analyses, including in the same analysis plots from all three habitat types. Plot species composition differences were analyzed using a non-metric multidimensional scaling (NMDS) analysis based on Bray–Curtis dissimilarities calculated on abundance data (expressed as percentages). We further evaluated the role of particular species in the species turnover due to the
invasion process by carrying out an Indicator Species Analysis (ISA, [56]). The ISA allows computing an indicator value \( d \) (ranging between 0 and 100) of each species as the product of the relative frequency and relative average abundance of species in clusters. The analysis also produces a significance value, representing the probability of obtaining a \( d \) value as high as that observed over 9999 iterations. We conducted the analyses by merging the two years of survey and adopting a grouping based on status and invasion and habitat.

Finally, we further evaluated the beta diversity patterns between invaded and control plots within each habitat type for both years, calculating the distance matrices accounting for spatial turnover, nestedness, and the sum of both components [37], using species presence/absence data. We used Sørensen’s index to quantify dissimilarity.

All analyses were conducted in R environment (R version 4.1.0): the GLS models were fitted using the ‘nlme’ package version 3.1-15 [57]; the NMDS was produced using the ‘vegan’ package version 2.5-7 [58]; the ISA was conducted using the package ‘labdsv’ (R package version 1.8-0, https://CRAN.R-project.org/package=labdsv, accessed on 16 October 2022) and the beta diversity was calculated using the package ‘betapart’ [59]. All plots were drawn using ‘ggplot2’ package version 3.3.3 [60].

**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/plants11202802/s1, Table S1: Environmental variables of the 88 sampled plots; Table S2: species per plot community matrix, with plots in the rows and species in the columns; Table S3: Vegetation total mean cover values of sampled plots; Table S4: abbreviations for plant names used in Table S2 and Figure 4; Table S5: Climatic data for Giglio Island.

**Author Contributions:** Conceptualization, L.L. and M.M.; methodology, L.L. and M.M.; formal analysis, L.L. and M.M.; field investigation, L.L., G.F. and M.M.; data curation, L.L. and M.M.; writing—original draft preparation, L.L. and M.M.; writing—review and editing, R.B., F.G., M.G., G.F., L.L., M.M. and D.V. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was realized within the LIFE Nature and Biodiversity project LIFE LETSGO GIGLIO “Less alien species in the Tuscan Archipelago: new actions to protect Giglio island habitats” (LIFE18 NAT/IT/000828).

**Institutional Review Board Statement:** Not applicable.

**Informed Consent Statement:** Not applicable.

**Data Availability Statement:** Data are contained within the article or Supplementary Material.

**Acknowledgments:** We thank Lorella Dell’Olmo for the realization of Figure 5. We wish to thank the other partners of the LIFE LETSGO GIGLIO project, the Tuscan Archipelago National Park and NEMO Nature and Environment Management Operators s.r.l., for their support.

**Conflicts of Interest:** The authors declare no conflict of interest.

**References**

1. Diaz, S.M.; Settele, J.; Brondizio, E.; Ngo, H.; Guéze, M.; Agard, J.; Arneth, A.; Balvanera, P.; Braunam, K.; Butchart, S.; et al. The Global Assessment Report on Biodiversity and Ecosystem Services: Summary for Policy Makers; Intergovernmental Science-Policy Platform on Biodiversity and Ecosystem Services secretariat: Bonn, Germany, 2019; ISBN 978-3-947851-13-3.

2. Pyšek, P.; Hulme, P.E.; Simberloff, D.; Bacher, S.; Blackburn, T.M.; Carlton, J.T.; Dawson, W.; Essl, F.; Foxcroft, L.C.; Genovesi, P.; et al. Scientists’ Warning on Invasive Alien Species. *Biol. Rev.* 2020, 95, 1511–1534. [CrossRef] [PubMed]

3. Seebens, H.; Blackburn, T.M.; Dyer, E.E.; Genovesi, P.; Hulme, P.E.; Jeschke, J.M.; Pagad, S.; Pyšek, P.; Winter, M.; Arianoutsou, M.; et al. No Saturation in the Accumulation of Alien Species Worldwide. *Nat. Commun.* 2017, 8, 14435. [CrossRef] [PubMed]

4. Hejda, M.; Pyšek, P.; Jarosňik, V. Impact of Invasive Plants on the Species Richness, Diversity and Composition of Invaded Communities. *J. Ecol.* 2009, 97, 393–403. [CrossRef]

5. Vilà, M.; Espinar, J.L.; Hejda, M.; Hulme, P.E.; Jarosňik, V.; Maron, J.L.; Pergl, J.; Schaffner, U.; Sun, Y.; Pyšek, P. Ecological Impacts of Invasive Alien Plants: A Meta-Analysis of Their Effects on Species, Communities and Ecosystems. *Ecol. Lett.* 2011, 14, 702–708. [CrossRef]

6. Vilà, M.; Rohr, R.P.; Espinar, J.L.; Hulme, P.E.; Pergl, J.; Le Roux, J.J.; Schaffner, U.; Pyšek, P. Explaining the Variation in Impacts of Non-Native Plants on Local-Scale Species Richness: The Role of Phylogenetic Relatedness. *Glob. Ecol. Biogeogr.* 2015, 24, 139–146. [CrossRef]
7. Vilà, M.; Hulme, P.E. Impact of Biological Invasions on Ecosystem Services; Springer: Cham, Switzerland, 2017.

8. Kumar Rai, P.; Singh, J.S. Invasive Alien Plant Species: Their Impact on Environment, Ecosystem Services and Human Health. Ecol. Indic. 2020, 111, 106020. [CrossRef]

9. Olden, J.D.; Rooney, T.P. On Defining and Quantifying Biotic Homogenization. Glob. Ecol. Biogeogr. 2006, 15, 113–120. [CrossRef]

10. Dawson, W.; Moser, D.; van Kleunen, M.; Kreft, H.; Pergl, J.; Pyšek, P.; Weigelt, P.; Winter, M.; Lenzner, B.; Blackburn, T.M.; et al. Global Hotspots and Correlates of Alien Species Richness across Taxonomic Groups. Nat. Ecol. Evol. 2017, 1, 0186. [CrossRef]

11. Gaertner, M.; Den Breejen, A.; Hui, C.; Richardson, D.M. Impacts of Alien Plant Invasions on Species Richness in Mediterranean-Type Ecosystems: A Meta-Analysis. Prog. Phys. Geogr. Earth Environ. 2009, 33, 319–338. [CrossRef]

12. Brundu, G. Invasive Alien Plants in Protected Areas in Mediterranean Islands: Knowledge Gaps and Main Threats. In Plant Invasions in Protected Areas: Patterns, Problems and Challenges; Foxcroft, L.C., Pyšek, P., Richardson, D.M., Genovesi, P., Eds.; Invading Nature—Springer Series in Invasion Ecology; Springer: Dordrecht, The Netherlands, 2013; pp. 395–422. ISBN 978-94-007-7750-7.

13. Celesti-Grapow, L.; Bassi, L.; Brundu, G.; Camarda, I.; Carli, E.; D’Auria, G.; Del Guacchio, E.; Domina, G.; Ferretti, G.; Foggi, B.; et al. Plant Invasions on Small Mediterranean Islands: An Overview. Plant Biosyst.-Int. J. Deal. Asp. Plant Biol. 2016, 150, 1119–1133. [CrossRef]

14. Lazzaro, L.; Foggi, B.; Ferretti, G.; Brundu, G. Priority Invasive Alien Plants in the Tuscan Archipelago (Italy): Comparing the EPPO Prioritization Scheme with the Australian WRA. Biol. Invasions 2016, 18, 1317–1333. [CrossRef]

15. Lazzaro, L.; Bolpagni, R.; Buffa, G.; Gentili, R.; Lonati, M.; Stinca, A.; Acosta, A.T.R.; Adorni, M.; Aleffi, M.; Allegrezza, M.; et al. Impact of Invasive Alien Plants on Native Plant Communities and Natura 2000 Habitats: State of the Art, Gap Analysis and Perspectives in Italy. J. Environ. Manag. 2020, 274, 111140. [CrossRef] [PubMed]

16. Maes, J. A Model for the Assessment of Habitat Conservation Status in the EU; Publication Office of the European Commission JRC scientific and policy report; European Commission Joint Research Centre Institute for Environment and Sustainability: Luxembourg, 2013.

17. Pyšek, P.; Jarosík, V.; Hulme, P.E.; Pergl, J.; Hejda, M.; Schaffner, U.; Vilà, M. A Global Assessment of Invasive Plant Impacts on Resident Species, Communities and Ecosystems: The Interaction of Impact Measures, Invading Species’ Traits and Environment. Glob. Chang. Biol. 2012, 18, 1725–1737. [CrossRef]

18. Guerra, C.; Baquero, R.A.; Gutiérrez-Arellano, D.; Nicola, G.G. Is the Natura 2000 Network Effective to Prevent the Biological Invasions? Glob. Ecol. Conserv. 2018, 16, e00497. [CrossRef]

19. Mazaris, A.D.; Katsanevakis, S. The Threat of Biological Invasions Is Under-Represented in the Marine Protected Areas of the European Natura 2000 Network. Biol. Conserv. 2018, 225, 208–212. [CrossRef]

20. Campoy, J.G.; Acosta, A.T.R.; Affre, L.; Barreiro, R.; Brundu, G.; Buisson, E.; González, L.; Lema, M.; Novoa, A.; Retuerto, R.; et al. Monographs of Invasive Plants in Europe: Carpobrotus. Bot. Lett. 2018, 165, 440–475. [CrossRef]

21. Carranza, M.L.; Carboni, M.; Feola, S.; Acosta, A.T.R. Landscape-Scale Patterns of Alien Plant Species on Coastal Dunes: The Case of Iceplant in Central Italy. Appl. Veg. Sci. 2010, 13, 135–145. [CrossRef]

22. Molinari, N.; D’Antonio, C.; Thomson, G.; Cuddington, K.; Byers, J.; Hastings, A.; Wilson, W. Carpobrotus as a Case Study of the Complexities of Species Impacts. In Ecosystem Engineers: Plants to Protists; Theroretical ecology series; Elsevier: London, UK, 2007; pp. 139–162.

23. Santoro, R.; Jucker, T.; Carboni, M.; Acosta, A.T.R. Patterns of Plant Community Assembly in Invaded and Non-Invaded Communities along a Natural Environmental Gradient. J. Veg. Sci. 2012, 23, 483–494. [CrossRef]

24. Vilà, M.; Tessier, M.; Suehs, C.M.; Brundu, G.; Carta, L.; Galanidis, A.; Lambdon, P.; Manca, M.; Médail, F.; Moragues, E.; et al. Local and Regional Assessments of the Impacts of Plant Invaders on Vegetation Structure and Soil Properties of Mediterranean Islands. J. Biogeogr. 2006, 33, 853–861. [CrossRef]

25. Acosta, A.; Izza, C.F.; Stanisci, A. Comparison of Native and Alien Plant Traits in Mediterranean Coastal Dunes. Community Ecol. 2006, 7, 35–41. [CrossRef]

26. Santoro, R.; Carboni, M.; Carranza, M.L.; Acosta, A.T.R. Focal Species Diversity Patterns Can Provide Diagnostic Information on Plant Invasions. J. Nat. Conserv. 2012, 20, 85–91. [CrossRef]

27. Jucker, T.; Carboni, M.; Acosta, A.T.R. Going beyond Taxonomic Diversity: Deconstructing Biodiversity Patterns Reveals the True Cost of Iceplant Invasion. Divers. Conserv. 2013, 19, 1566–1577. [CrossRef]

28. Badalamenti, E.; Cristina, L.; Laudicina, V.A.; Novara, A.; Pasta, S.; La Mantia, T. The Impact of Carpobrotus Cfr. Acinaciformis (L.) Bolus on Soil Nutrients, Microbial Communities Structure and Native Plant Communities in Mediterranean Ecosystems. Plant Soil 2016, 409, 19–34. [CrossRef]

29. Santoro, R.; Jucker, T.; Carranza, M.; Acosta, A. Assessing the Effects of Carpobrotus Invasion on Coastal Dune Soils. Does the Nature of the Invaded Habitat Matter? Community Ecol. 2011, 12, 234–240. [CrossRef]

30. Novoa, A.; González, L.; Moravcová, L.; Pyšek, P. Constraints to Native Plant Species Establishment in Coastal Dune Communities Invaded by Carpobrotus Edulis: Implications for Restoration. Biol. Conserv. 2013, 164, 1–9. [CrossRef]

31. Vieites-Blanco, C.; González-Prieto, S.J. Effects of Carpobrotus Edulis Invasion on Soil Gross N Fluxes in Rocky Coastal Habitats. Sci. Total Environ. 2018, 619–620, 966–976. [CrossRef] [PubMed]

32. Novoa, A.; González, L. Impacts of Carpobrotus Edulis (L.) N.E.Br. on the Germination, Establishment and Survival of Native Plants: A Clue for Assessing Its Competitive Strength. PLoS ONE 2014, 9, e107557. [CrossRef]
33. Novoa, A.; González, L.; Moravcová, L.; Pyšek, P. Effects of Soil Characteristics, Allelopathy and Frugivory on Establishment of the Invasive Plant Carpobrotus Edulis and a Co-Ocurring Native, Malcolmia Littorea. *PLoS ONE* 2012, 7, e3166. [CrossRef]

34. Brunel, S.; Brundu, G.; Fried, G. Eradication and control of invasive alien plants in the Mediterranean Basin: Towards better coordination to enhance existing initiatives. *EPPO Bull.* 2013, 43, 290–308. [CrossRef]

35. Lazzaro, L.; Ferretti, G.; Giuliani, C.; Foggì, B. A Checklist of the Alien Flora of the Tuscan Archipelago (Italy). *Webbia* 2014, 69, 157–176. [CrossRef]

36. Hulme, P.E.; Pyšek, P.; Jarosík, V.; Pergl, J.; Schaffner, U.; Vilà, M. Bias and Error in Understanding Plant Invasion Impacts. *Trends Ecol. Evol.* 2013, 28, 212–218. [CrossRef] [PubMed]

37. Baselga, A. Partitioning the Turnover and Nestedness Components of Beta Diversity. *Glob. Ecol. Biogeogr.* 2010, 19, 134–143. [CrossRef]

38. Chase, J.M.; Myers, J.A. Disentangling the Importance of Ecological Niches from Stochastic Processes across Scales. *Philos. Trans. R. Soc. B Biol. Sci.* 2011, 366, 2351–2363. [CrossRef] [PubMed]

39. Leprieur, F.; Olden, J.D.; Lek, S.; Brosse, S. Contrasting Patterns and Mechanisms of Spatial Turnover for Native and Exotic Freshwater Fish in Europe. *J. Biogeogr.* 2009, 36, 1899–1912. [CrossRef]

40. Marini, L.; Bertolli, A.; Bona, E.; Federici, G.; Martini, F.; Prosser, F.; Bommarco, R. Beta-Diversity Patterns Elucidate Mechanisms of Alien Plant Invasion in Mountains. *Glob. Ecol. Biogeogr.* 2013, 22, 450–460. [CrossRef]

41. Fried, G.; Laitung, B.; Pierre, C.; Chagu, ë.

42. Marzialetti, F.; Cascone, S.; Frate, L.; Di Febbraro, M.; Acosta, A.T.R.; Carranza, M.L. Measuring Alpha and Beta Diversity by reed canary grass (*Phalaris arundinacea*). *Biol. Invasions 2010*, 12, 1253–1267. [CrossRef]

43. Buisson, E.; Braschi, J.; Chenet-Lescure, J.; Hess, M.C.M.; Vidalier, C.; Pavon, D.; Ramone, H.; Amy-Krebs, E.; Cottaz, C.; Pasetti, A. Native Plant Community Recovery after Carpobrotus (Ice Plant) Removal on an Island—Results of a 10-Year Project. *Appl. Veg. Sci.* 2021, 24, e12524. [CrossRef]

44. Foggì, B.; Fancioli, V. Contribution to the Knowledge of the Vegetation of the Giglio Island (Tuscan Archipelago, Southern Tuscany). *Webbia* 2008, 63, 25–48. [CrossRef]

45. Conser, C.; Connor, E.F. Assessing the Residual Effects of Carpobrotus Edulis Invasion, Implications for Restoration. *Biol. Invasions 2009*, 11, 349–358. [CrossRef]

46. Malavasi, M.; Santoro, R.; Cutini, M.; Acosta, A.T.R.; Carranza, M.L. The Impact of Human Pressure on Landscape Patterns and Plant Species Richness in Mediterranean Coastal Dunes. *Plant Biosyst.-Int. J. Deel. Asp. Plant Biol.* 2016, 150, 73–82. [CrossRef]

47. Spyreas, G.; Wilm, B.W.; Plocher, A.E.; Ketzenr, D.M.; Matthews, J.W.; Ellis, J.L.; Heske, E.J. Biological consequences of invasion by reed canary grass (*Phalaris arundinacea*). *Biol. Invasions 2010*, 12, 1253–1267. [CrossRef]

48. Benesperi, R.; Giuliani, C.; Zanetti, S.; Gennai, M.; Mariotti Lippi, M.; Guidi, T.; Nascimbene, J.; Foggì, B. Forest plant diversity is threatened by *Robinia pseudoacacia* (black-locust) invasion. *Biodivers. Conserv.* 2012, 21, 3555–3568. [CrossRef]

49. Chen, G.Q.; Zhang, C.B.; Ling, M.A.; Qiang, S.; Silander, J.A.; Qi, L.L. Biotic homogenization caused by the invasion of reed canary grass (*Phalaris arundinacea*). *Biol. Invasions 2010*, 12, 1253–1267. [CrossRef]

50. Malavasi, M.; Santoro, R.; Cutini, M.; Acosta, A.T.R.; Carranza, M.L. The Impact of Human Pressure on Landscape Patterns and Plant Species Richness in Mediterranean Coastal Dunes. *Plant Biosyst.-Int. J. Deel. Asp. Plant Biol.* 2016, 150, 73–82. [CrossRef]

51. Spyreas, G.; Wilm, B.W.; Plocher, A.E.; Ketzenr, D.M.; Matthews, J.W.; Ellis, J.L.; Heske, E.J. Biological consequences of invasion by reed canary grass (*Phalaris arundinacea*). *Biol. Invasions 2010*, 12, 1253–1267. [CrossRef]

52. Dyderski, M.K.; Jagodziński, A.M. Impacts of Invasive Trees on Alpha and Beta Diversity of Temperate Forest Understories. *Biol. Invasions* 2021, 23, 235–252. [CrossRef]

53. Marzialetti, F.; Cascone, S.; Frate, L.; Di Febbraro, M.; Acosta, A.T.R.; Carranza, M.L. Measuring Alpha and Beta Diversity by Field and Remote-Sensing Data: A Challenge for Coastal Dunes Biodiversity Monitoring. *Remote Sens.* 2021, 13, 1928. [CrossRef]

54. Baldini, R.M. The Vascular Flora of the Giglio Island. (Tuscan Archipelago): Taxonomic Revision and Updating. *Webbia* 1998, 52, 307–404. [CrossRef]

55. Foggì, B.; Viciani, D.; Baldini, R.M.; Carta, A.; Guidi, T. Conservation Assessment of the Endemic Plants of the Tuscan Archipelago, Italy. *Orx. 2015*, 49, 118–126. [CrossRef]

56. Dufrené, M.; Legendre, P. Species Assemblages and Indicator Species: The Need for a Flexible Asymmetrical Approach. *Ecol. Monogr.* 1997, 67, 345–366. [CrossRef]

57. Pinheiro, J.; Bates, D. *Mixed-Effects Models in S and S-PLUS*; Springer Science & Business Media: Berlin/Heidelberg, Germany, 2006; p. 528.

58. Oksanen, J.; Blanchet, F.G.; Friendly, M.; Kindt, R.; Legendre, P.; McGlinn, D.; Minchin, P.R.; O’Hara, R.B.; Simpson, G.L.; Solymos, P.; et al. Vgenc: *Community Ecology Package*, 2020.

59. Baselga, A.; Orme, D.; Villeger, S.; De Bortoli, J.; Leprieur, F.; Baselga, M.A. Package ‘betapart’. Partitioning beta diversity into turnover and nestedness components, version 1.5.6. 2022.

60. Wickham, H. *Ggplot2: Elegant Graphics for Data Analysis*; Springer: New York, NY, USA, 2016.