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## Impacts of alien plant invasions on species richness in Mediterranean-type ecosystems: a meta-analysis

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Abstract: Besides a general consensus regarding the negative impact of invasive alien species in the literature, only recently has the decline of native species attributable to biological invasions begun to be quantified in many parts of the world. The cause-effect relationship between the establishment and proliferation of alien species and the extinction of native species is, however, seldom demonstrated. We conducted a meta-analysis of studies in Mediterranean-type ecosystems (MTEs) to examine: (1) whether invasion of alien plant species indeed causes a reduction in the number of native plant species at different spatial and temporal scales; (2) which growth forms, habitat types and areas are most affected by invasions; and (3) which taxa are most responsible for native species richness declines. Our results confirm a significant decline in native species richness attributable to alien invasions. Studies conducted at small scales or sampled over long periods reveal stronger impacts of alien invasion than those at large spatial scales and over short periods. Alien species from regions with similar climates have much stronger impacts, with the native species richness in South Africa and Australia declining significantly more post-invasion than for European sites. Australian Acacia species in South Africa accounted for the most significant declines in native species richness. Among the different growth forms of alien plants, annual herbs, trees and creepers had the greatest impact, whereas graminoids generally caused insignificant changes to the native community. Native species richness of shrublands, old fields and dune vegetation showed significant declines, in contrast to insignificant declines for forest habitats.

**Key words:** biodiversity, biological invasions, exotic species, growth form, habitat type, spatial and temporal scale.

#### I Introduction

Invasive alien species are considered a threat to biodiversity and ecosystem stability, and

are widely held to be responsible for the decline of native species richness and the local extinction of certain species (Richardson

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et al., 1989; Wilcove et al., 1998; Davis, 2003). However, it is often difficult to demonstrate the cause-effect relationship between the establishment and proliferation of alien plant species and the extinction of native species (Brown and Sax, 2004; Davis, 2009). Recently, the broadly accepted connection between invasive species and native diversity declines has been debated (Houlahan and Findlay, 2004; Hejda and Pyšek, 2006; Richardson et al., 2007). Perceptions on the negative impacts of invasions on native plant communities are, to a certain degree, affected by the spatial scales of studies. This has been widely recognized in the discussion about the 'invasion paradox' which describes the co-occurrence of independent lines of support for both a negative and a positive relationship between native biodiversity and the invasions of invasive alien species depending on the spatial scale of investigation (Fridley et al., 2007). Another factor which must be considered when evaluating impacts of alien invasions on native species richness is the different timescales involved in invasions and extinctions (Sax et al., 2002; Richardson et al., 2007). Although some species may be doomed to extinction due to disruptions caused by invasions (eg, Traveset and Richardson, 2006), extinction can be delayed (Tilman et al., 1994). With plant species especially, the process of extinction takes much longer (decades or longer) than naturalizations or invasion (rates easily measured in years). It is likely that many plant species will eventually be driven to extinction as a direct consequence of current processes. Therefore, we would expect a stronger signal of negative impact of invasions from studies that capture effects over longer periods. It is also important to consider that although invasions of alien species may not result in extinctions of entire species (over the typically short timescale captured in a field study) this does not mean that they are not reducing biodiversity. They may still be causing declines in the abundance of native species or the elimination of some populations which

may reduce genetic diversity (Davis, 2009). Nevertheless, the above generalizations have emerged from isolated case studies and it is difficult to synthesize results from different spatial and temporal scales. Besides these scale-effects on the observed impact of alien invasions, features of the invading species and the invaded habitat must also be taken into account when considering the impact of alien invasions on native species richness.

We investigated how different spatial and temporal scales and factors (such as taxa, growth forms, habitat types and countries) affect the impact of invasions on native species richness. We chose to study the impact of invasions on native species richness in Mediterranean-type ecosystems (MTEs) which have similar climates and other environmental drivers. Several studies have addressed the importance of comparing alien plant invasions in similar climatic regions (Kruger et al., 1989; Sax, 2002; Pauchard et al., 2004), as comparing ecosystems with widely diverging climates, disturbance regimes and other factors may reduce our ability to isolate the effects of particular stressors. Mediterranean-climate zones are considered to be especially appropriate for global-scale 'natural experiments' as they differ less in key aspects than other biome types that occur at multiple localities around the world, eg, savannas (Pauchard et al., 2004). Invasions in MTEs have been well studied for decades (Groves and Di Castri, 1991; Richardson et al., 1992; Rejmánek and Randall, 1994) and the problem of alien plant invasions is widely recognized as a major threat to biodiversity in all MTEs today (Rejmánek and Randall, 1994; Rouget et al., 2003; Seabloom et al., 2006; Underwood et al., 2009).

We conducted a meta-analysis to quantify the impact of alien invasion on native plant species richness. This approach allowed us to examine: (1) whether invasion of alien plant species indeed causes a reduction in the number of native plant species at different localities and at different spatial and temporal scales; (2) which growth forms, habitat types and areas are most affected by invasions; and (3) which taxa are most responsible for native species richness declines.

#### **II Methods**

#### 1 Mediterranean-type ecosystems

Mediterranean-type ecosystems (MTEs) are found between latitudes 32° and 40° north and south of the equator on the west coast of continents in five regions of the world, namely South Africa, Australia, California, Chile and the Mediterranean Basin (Aschmann, 1973). The defining factors for the Mediterraneantype climate are summer drought and winter rainfall (Köppen, 1923). Annual rainfall ranges from below 90 mm to 1500 mm; annual mean temperature ranges from about 11°C to 17°C; mean seasonal temperatures are 8°C in the coldest month and up to 25°C in the hottest month (Aschmann, 1973). The vegetation of MTEs is characterized by chaparral-like shrublands, coastal scrub, woodland and forest (Dallman, 1998). MTEs have remarkable plant diversity. Covering less than 5% of the Earth's surface, they contain nearly 20% of the planet's known plant species (Cowling et al., 1996).

#### 2 Data

We confined our search to MTEs in five regions covering South Africa, Australia, Europe, California and Chile. The first criterion for our literature search was that the focus of the study should be an invasive plant species alien to the area under investigation (sensu Richardson et al., 2000; sensu Pyšek et al., 2004). Second, we only included studies that directly compared invaded ecosystems dominated by invasive alien species with corresponding, relatively intact, ecosystems in terms of native species richness. Third, we also required that species richness was quantified (or where this could be computed from presented data) as mean number of native species with standard error (SE) and sample size. In some cases, where such data were not provided, we obtained the data from the authors.

We searched for papers using combinations of the terms 'exotic', 'invasive', 'invasion', 'alien species', 'species richness' and 'biodiversity' on 'Web of Science', 'JSTOR' and 'Google Scholar'. Additional literature was obtained through conventional searches of the bibliographies of papers and reports. We did not limit the review to papers in a set of journals published during a certain period. This method allowed us to include a wider range of literature than if we had limited the survey to only certain journals or a certain timeframe.

#### 3 Analysis

We summarized data from all the studies that fulfilled our criteria in a table including information about study area and habitat, origin and growth form of the invading species, and temporal and spatial scale, defined by extent of study area and unit size (grain) (Table 1). Eleven studies (three from South Africa, four from Europe, three from California and one from Australia) including 24 species (Figure 1) met our criteria. Some studies investigated species at different sites and in different seasons this left us with 47 cases for the meta-analysis. Chile was not included in the meta-analysis at all as none of the studies met our criteria.

For the meta-analysis, we recorded the number of native species (mean and SE) in invaded and natural reference sites for each study. The meta-analysis is a technique of quantitative research synthesis (Smith and Glass, 1977) and has been widely used in ecology (eg, Ashton *et al.*, 2005). In this study, we used the comprehensive meta-analysis software (CMA version 2.0; Borenstein *et al.*, 2005) to conduct a twogroup comparison (native versus invaded) with additional moderators as defined by, for example, taxa, temporal scale, spatial scale, growth form, habitat, origin or invaded country. Cohen's (1988) mean difference

spec	species richness in Mediterranean-type ecosystems	1editerranean-	type ecosys	tems				·	·
Case	Species	Country of origin	Invaded	Timescale <sup>4</sup>	Unit size	Extent (km²)	Growth form	Habitat	Reference
		:			()				
_	Acacia saligna	Australia	South Africa	S	_	0.00012	tree	shrubland	Holmes and Cowling, 1997
2	Acacia saligna	Australia	South Africa	s	_	0.00012	tree	shrubland	Holmes and Cowling, 1997
З	Acacia saligna	Australia	South Africa	S	1	0.00012	tree	shrubland	Holmes and Cowling, 1997
4	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
5	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
9	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
2	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
∞	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
6	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
10	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
Ξ	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante et al., 2003
12	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
13	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
14	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
15	Acacia longifolia	Australia	Portugal	S	25	0.003	tree	dune vegetation	Marchante <i>et al.</i> , 2003
16	Acacia melanoxylon	Australia	South Africa	S	4	0.01	tree	shrubland	Richardson <i>et al.</i> , 1989
17	Pinus pinaster	Europe	South Africa	S	4	0.01	tree	shrubland	Richardson <i>et al.</i> , 1989
18	different species <sup>1</sup>	South Africa/ Australia/Europe	South Africa	S	4	0.01	tree	different types	Richardson <i>et al.</i> , 1989
19	Asparagus asparagoides	South Africa	Australia	S	10	0.0004	creeper	shrubland	Turner et al., 2008
20	Ailanthus altissima	China	Europe	S	4	302	tree	different types	Vilà <i>et al.</i> , 2006
21	Ailanthus altissima	China	Europe	S	4	3656	tree	different types	Vilà <i>et al.</i> , 2006
22	Ailanthus altissima	China	Europe	S	4	8682	tree	different types	Vilà <i>et al.</i> , 2006
23	Ailanthus altissima	China	Europe	S	4	24090	tree	different types	Vilà <i>et al.</i> , 2006
24	Ailanthus altissima	China	Europe	S	4	8700	tree	different types	Vilà <i>et al.</i> , 2006
25	Carpobrotus spp.	South Africa	Europe	S	4	3656	succulent	different types	Vilà <i>et al.</i> , 2006

 Table 1a
 Information on studies used in the meta-analysis to determine impacts of invasive plants on native plant

26	Carpobrotus spp.	South Africa	Europe	S	4	302	succulent	different types	Vilà <i>et al.</i> , 2006
27	Carpobrotus spp.	South Africa	Europe	S	4	6	succulent	different types	Vilà <i>et al.</i> , 2006
28	Carpobrotus spp.	South Africa	Europe	S	4	12	succulent	different types	Vilà <i>et al.</i> , 2006
29	Carpobrotus spp.	South Africa	Europe	S	4	24090	succulent	different types	Vilà <i>et al.</i> , 2006
30	Carpobrotus spp.	South Africa	Europe	S	4	8700	succulent	different types	Vilà <i>et al.</i> , 2006
31	Oxalis pes-caprae	South Africa	Europe	S	4	3656	perennial herb	different types	Vilà <i>et al.</i> , 2006
32	Oxalis pes-caprae	South Africa	Europe	S	4	302	perennial herb	different types	Vilà <i>et al.</i> , 2006
33	Oxalis pes-caprae	South Africa	Europe	S	4	1632		different types	Vilà <i>et al.</i> , 2006
34	Oxalis pes-caprae	South Africa	Europe	S	4	12	succulent	different types	Vilà <i>et al.</i> , 2006
35	Oxalis pes-caprae	South Africa	Europe	S	4	24090	perennial herb	different types	Vilà <i>et al.</i> , 2006
36	Oxalis pes-caprae	South Africa	Europe	S	4	8700	perennial herb	different types	Vilà <i>et al.</i> , 2006
37	Cortaderia jubata	Southern America California	a California	S	2500	0.025	graminoid	shrubland	Lambrinos, 2000
38	Cortaderia jubata	Southern America California	a California	S	2500	0.025	graminoid	shrubland	Lambrinos, 2000
39	Cortaderia jubata	Southern America California	a California	S	2500	0.025	graminoid	shrubland	Lambrinos, 2000
40	Cortaderia jubata	Southern America California	a California	S	2500	0.025	graminoid	shrubland	Lambrinos, 2000
41	Delairea odorata	South Africa	California	S	2,25	-	lerb	shrubland	Alvarez and Cushman, 2002
42	Delairea odorata	South Africa	California	S	2,25	-	perennial herb shrubland	shrubland	Alvarez and Cushman, 2002
43	Delairea odorata	South Africa	California	S	2,25	_	perennial herb shrubland	shrubland	Alvarez and Cushman, 2002
44	Fallopia japonica	Europe	Belgium	S	6	10	perennial herb forest	forest	Vanderhoeven et al., 2005
45	different species <sup>2</sup>	California	California	_	10	0.01	annual herb	shrubland	Stylinski and Allen, 1999
46	Oxalis pes-caprae	Greece/South Africa	Europe	S	0,25	0.004	perennial herb	old field	Petsikos <i>et al.</i> , 2007
47	exotic annual species <sup>3</sup>	South Africa/ Europe	South Africa	S	_	625	annual herb	shrubland	Vlok, 1988
<sup>1</sup> Pinu <sup>2</sup> Hvp	<sup>1</sup> Pinus spp., P. pinaster, P. radiata, Hakea sericea, Acacia sali <sup>2</sup> Hvpochaeris glabra. Bromus spp., Erodium botry, Avena spp.	<sup>D</sup> . radiata, Hakea : mus spp., Erodiun	Hakea sericea, Acacia saligna, Acacia melanoxylon Erodium botrv. Avena spp.	saligna, Aco spp.	acia melano	nolyx			

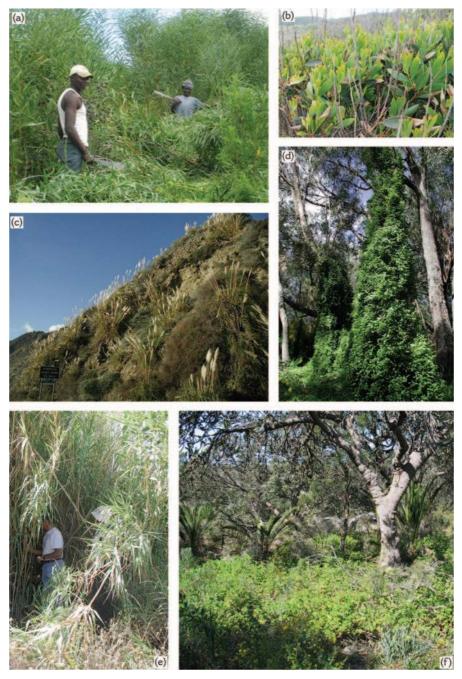
<sup>2</sup>Hypochaeris glabra, Bromus spp., Erodium botry, Avena spp.
<sup>3</sup>Bromus diandrus, Hordeum murinum ssp. glaucum, Lolium perenne, Lolium rigidum, Lolium temulentum, Phalaris canariensis, Vulpia bromoides
<sup>4</sup>s = short-term study (2–3 yrs); l = long-term study (>10 yrs)

Deti	Details for case studies are given in Table la	are given in T	able la		•		~		
		Effect size and 95% confidence interval	5% confidence int	erval		Test of null (2-tail)	tail)		
Case study	Case Species study	رم ه	SE <sup>b</sup>	Lower CI95°	Upper CI95 <sup>d</sup>	Z–value <sup>e</sup>	p-value	Weight	Residual
_	Acacia saligna	-4.627734781	0.428776816	-5.4681219	-3.7873477	-10.792875	0	2.245591114	-3.1245763
2	Acacia saligna	-5.699759745	0.503036164	-6.6856925	-4.713827	-11.330716	0	2.13363434	-3.9637791
3	Acacia saligna	-3.106208205	0.33211942	-3.7571503	-2.4552661	-9.3526847	0	2.378228061	-1.8376697
4	Acacia longifolia	-0.787238207	0.599296254	-1.9618373	0.38736087	-1.3136044	0.188979361	1.982095738	0.2420421
2	Acacia longifolia	-1.082990516	0.618225568	-2.2946904	0.12870933	-1.7517724	7.98E-02	1.951984239	-2.34E-03
6	Acacia longifolia	-0.723057872	0.595917104	-1.8910339	0.44491819	-1.2133531	0.224994845	1.987468499	0.2954859
7	Acacia longifolia	-0.877865514	0.604519364	-2.0627017	0.30697067	-1.452171	0.146454052	1.973788928	0.1667936
∞	Acacia longifolia	-0.487264088	0.585855012	-1.6355188	0.66099064	-0.8317145	0.405570134	2.003457533	0.4926155
6	Acacia longifolia	-1.598928254	0.663217271	-2.8988102	-0.2990463	-2.4108664	1.59E-02	1.880517309	-0.4173606
10	Acacia longifolia	-0.282437234	0.58022161	-1.4196507	0.85477622	-0.4867748	0.62641796	2.012401134	0.6643126
11	Acacia longifolia	-0.996434003	0.612130185	-2.1961871	0.20331911	-1.6278139	0.103564359	1.961681662	6.88E-02
12	Acacia longifolia	9.92E-02	0.577705547	-1.0330411	1.23152306	0.1717847	0.863606764	2.016393334	0.9831613
13	Acacia longifolia	-0.669646514	0.593310854	-1.8325144	0.49322139	-1.1286605	0.259041086	1.991611414	0.340045
14	Acacia longifolia	0.123452682	0.577899954	-1.0092104	1.25611578	0.2136229	0.830841116	2.016084926	1.0032665
15	Acacia longifolia	-2.12E-02	0.57736652	-1.1528391	1.11039606	-3.68E-02	0.970679778	2.016931145	0.8828609
16	Acacia melanoxylon	-7.689849727	1.832127026	-11.280753	-4.0989467	-4.1972252	2.70E-05	0.65930125	-3.1291088
17	Pinus pinaster	-0.730112887	0.540120758	-1.7887301	0.32850435	-1.3517586	0.176452548	2.075776324	0.2961463
18	different species <sup>1</sup>	-2.391637231	0.826301885	-4.0111592	-0.7721153	-2.8943868	3.80E-03	1.629195873	-0.9808011
19	Asparagus asparagoides	-2.11805426	0.558707299	-3.2131004	-1.0230081	-3.7909909	1.50E-04	2.046478466	-0.8717965
20	Ailanthus altissima	-0.551552832	0.330553971	-1.1994267	9.63E-02	-1.6685712	9.52E-02	2.380211081	0.4796395
21	Ailanthus altissima	-0.510541237	0.299649294	-1.0978431	0.07676059	-1.7037959	8.84E-02	2.418064758	0.5210482
22	Ailanthus altissima	-0.405798429	0.274952393	-0.9446952	0.13309836	-1.4758862	0.139974458	2.44641305	0.6205587
23	Ailanthus altissima	0.16	0.283294899	-0.3952478	0.7152478	0.5647825	0.572221715	2.437036677	1.1389876

**Table 1b** Effect sizes of all 47 studies. g = Cohen's (1988) mean difference effect size; SE = standard error; Lower C195 = lower confidence interval (95%); Upper C195 = upper confidence interval (95%); Z value = two-tail Z-test.

74	Ailanthus altissima	10 233020601	0 400631406	_1 0358835	0 56984211	70 5688546	0 569454848	0 073061814	0 7507966
1									
25	Carpobrotus spp.	-0.951036586	0.298403596	-l.5358969	-0.3661763	-3.1870815	1.44E-03	2.419536639	0.1181356
26	Carpobrotus spp.	-1.108831906	0.310065655	-1.7165494	-0.5011144	-3.5761197	3.49E-04	2.405589427	-2.62E-02
27	Carpobrotus spp.	-0.753789476	0.405922467	-1.5493829	4.18E-02	-1.8569789	6.33E-02	2.278552727	0.2895865
28	Carpobrotus spp.	-0.430497602	0.312160829	-1.0423216	0.18132638	-1.3790891	0.167867279	2.403044521	0.592375
29	Carpobrotus spp.	-0.877876225	0.270349568	-1.4077516	-0.3480008	-3.2471893	1.17E-03	2.45149631	0.1863298
30	Carpobrotus spp.	-1.802995743	0.448226538	-2.6815036	-0.9244879	-4.02251	5.76E-05	2.216915397	-0.6324912
31	Oxalis pes-caprae	3.08E-02	0.301529268	-0.5601461	0.62182686	0.1022798	0.918534583	2.415835244	1.0158044
32	Oxalis pes-caprae	-0.574818031	0.416593721	-1.3913267	0.24169066	-1.3798048	0.167646743	2.263267402	0.4468551
33	Oxalis pes-caprae	-0.739783497	0.266884488	-1.2628675	-0.2166995	-2.7719239	5.57E-03	2.455280116	0.3137921
34	Oxalis pes-caprae	-0.197900922	0.28938089	-0.765077	0.3692752	-0.683877	0.494052868	2.430066559	0.8090904
35	Oxalis pes-caprae	-4.07E-02	0.258225636	-0.5468254	0.46540052	-0.1576622	0.874722968	2.464571341	0.9601597
36	Oxalis pes-caprae	-1.613956446	0.514899393	-2.6231407	-0.6047722	-3.1345083	1.72E-03	2.115229984	-0.456009
37	Cortaderia jubata	-3.291858008	0.970472286	-5.1939487	-1.3897673	-3.3920165	6.94E-04	1.425972414	-1.5458383
38	Cortaderia jubata	-0.61588591	0.647275654	-1.8845229	0.65275106	-0.9515048	0.341348169	1.905801075	0.3760385
39	Cortaderia jubata	1.065397325	0.675835463	-0.2592158	2.39001049	1.5764152	0.114930138	1.860553922	1.72E+00
40	Cortaderia jubata	0.884877794	0.662684265	-0.4139595	2.18371509	1.3352932	0.181780403	1.881361639	1.5811996
41	Delairea odorata	-1.014466119	0.475108975	-1.9456626	-8.33E-02	-2.1352283	3.27E-02	2.176471766	5.69E-02
42	Delairea odorata	-0.900591374	0.469336362	-1.8204737	1.93E-02	-1.9188613	5.50E-02	2.185228031	0.1559506
43	Delairea odorata	-0.766964989	0.463363661	-1.6751411	0.1412111	-1.6552118	9.79E-02	2.194248208	0.2725882
44	Fallopia japonica	-0.684650429	0.514439811	-1.6929339	0.32363307	-1.330866	0.183233122	2.11594498	0.3379019
45	different species <sup>2</sup>	-2.997706948	0.505159926	-3.9878022	-2.0076117	-5.9341741	2.95E-09	2.130347783	-1.6440377
46	Oxalis pes-caprae	-1.832082371	0.284812292	-2.3903042	-1.2738605	-6.4325959	1.25E-10	2.435308994	-0.6903597
47	exotic annual species <sup>3</sup>	-1.637793969	0.270102937	-2.167186	-1.1084019	-6.0635918	1.33E-09	2.451766852	-0.5137536
		-1.080139574	0.172083842	-1.4174177	-0.7428614	-6.2768216	3.46E-10		
Dinus	Pinus spp., P. pinaster, P. radiata, Hakea sericea, Acacia saligna, Acacia melanoxylon	a, Hakea sericea, A	cacia saligna, Aca	cia melanoxylon					

Trius spp., г. рпизке, г. гишии, гимки зеткей, личии запуци, гичич телимулот. ²Hypochaeris glabra, Bromus spp., Erodium botry, Avena spp. ³Bromus diandrus, Hordeum murinum ssp. glaucum, Lolium perenne, Lolium rigidum, Lolium temulentum, Phalaris canariensis, Vulpia bromoides



**Figure 1** The impacts of invasive alien plants on native plant diversity have been investigated in many Mediterranean-type ecosystems around the world. Examples are: (a) *Acacia saligna* in South Africa; (b) *Acacia longifolia* in Portugal; (c) *Cortaderia jubata* in California, USA; (d) *Asparagus asparagoides* in Western Australia; (e) *Arundo donax* in California, USA; (f) *Pelargonium capitatum* in Western Australia (photo credits: (a) M. Gaertner; (b–f) D.M. Richardson)

effect size, g (Lipsey and Wilson, 2001; Borenstein *et al.*, 2005), and a mixed (random) effects model was used. A twotail Z-test was used to examine the null hypothesis (ie, the effect size equals zero) and a Q-test was used for the heterogeneity analysis. Meta-analysis can largely alleviate the bias of favouring significant results in literature by weighing each case according to its sampling variance and size, and, as a result, can present a more robust synthesis than traditional literature review.

#### **III Results**

The meta-analysis revealed an overall significant decline of species richness after invasion. In five of the 47 cases invasion had a positive effect on native species richness (species richness increased after invasion). In the remaining 42, invaded sites had lower native species richness. Eighteen cases (38.3%) had a negative effect size, indicating a significant decline of species richness after invasion, while 29 cases (61.7%) had effect sizes not significantly different from zero (ie, 95% CI includes 0), indicating no significant decline of species richness after invasion (Figure 2; Table 2). According to Cohen's (1988) standard, 48.9% of the cases had large effect sizes (>0.8), 36.2% had medium effect sizes  $(0.2 \sim 0.8)$ , and 14.9% had small effect sizes.

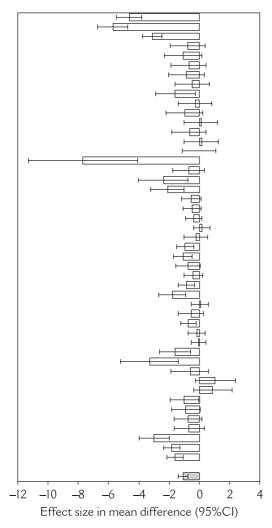
Effect size of invasion on native species richness declined significantly with increasing unit size. Cases with unit size <1 m<sup>2</sup>, 1–10 m<sup>2</sup> and 10–100 m<sup>2</sup> had effect sizes significantly different from zero (Z= -4.78; p < 0.001; Z= -5.97; p < 0.001; Z= -3.33; p < 0.001, respectively). Cases with unit sizes >100 m<sup>2</sup> had effect sizes not significantly different from zero (Z = -0.45; p = 0.65) (Figure 3a). The same pattern was evident for increasing extent of study area. Effect size was highest for case studies where the extent of the study area was less than 0.01 km<sup>2</sup>, and decreased with increasing extent of study area, except for studies with areal extent of 100-10 000 km<sup>2</sup>. Cases with study area extent <0.01 km<sup>2</sup>,  $1-100 \text{ km}^2$  and  $100-10\ 000 \text{ km}^2$  had effect sizes significantly different from zero (Z = -4.61; p < 0.001; Z = -2.46; p = 0.014; Z = -5;p < 0.001, respectively), whereas cases with extent of study area 0.01-1 km<sup>2</sup> and >10000km<sup>2</sup> had effect sizes not significantly different from zero (Z = -1.43; p = 0.153; Z = -0.81; p = 0.419, respectively) (Figure 3b). The time of investigation (temporal scale) also had a significant influence on effect size. Longterm investigations showed a significant higher effect size, with plant invasions having a stronger impact on native species richness than short-term investigations (Q = 13.51; p < 0.001).

Shrublands, old fields and dune vegetation showed significantly different effect sizes between groups (Q = 24.31; p < 0.001) with shrublands and old fields having largest declines in species richness attributable to alien invasion. Invaded sites in forest habitats. in contrast, showed no significant declines in species richness (Z = -1.33; p = 0.18). Among the different growth forms of alien plants reviewed, annual herbs, trees and creepers had the greatest impact on species richness decline (Z = -3.33; p < 0.001; Z = -3.78; p < 0.001; Z = -3.97; p < 0.001, respectively), whereas graminoids generally caused insignificant damage to the native communities (Z = -0.45; p = 0.65) (Figure 4a). An investigation of the different taxa included in the meta-analysis showed the highest effect size of invasion on species richness for two Australian Acacia species (A. melanoxylon and A. saligna) (Figure 4b; Table 2). Among the different countries investigated in the meta-analysis, South Africa had the highest declines of native plant species richness due to alien invasion followed by Australia (Figure 5a). Species with origin in Australia and Europe caused the largest declines in plant species richness (Figure 5b).

<b>Table 2</b> Summary of the Upper Cl95 = upper confidence	Summary of the = upper confidence	meta-analysis. SE = standard error; Lower interval (95%); Z value = two- tail Z-test; Q-value =	sis. SE = st ; Z value = tv	andard er vo- tail Z-te		C195 = lower confid Q-test for heterogeneity	er confic erogeneity	= lower confidence interval $(95\%)$ ; for heterogeneity	erval (	95%);
Model	Case study	Effect size		95% confidence interval	e interval	Test of null (2-tail)	(1)	Heterogeneity	ty	
		Point estimate	SEa	Lower CI95 <sup>b</sup>	Upper CI95°	Z-value <sup>d</sup>	p-value	Q-value <sup>e</sup>	df (Q)	p-value
Random effects	47	-1.080139574	0.172083842	-1.417418	-0.742861441	-6.276821588	3.5E-10			
Mixed effects analysis of unit size $(m^2)$	unit size (m²)									
- V	2ı	-3.337534451	0.697625712	-4.704856	-1.970213181	-4.784133373	1.7E-06			
1~10	26	-0.825721512	0.138366075	-1.096914	-0.55452899	-5.967658728	2.4E-09			
10~100 >100	12 4	-0.574693192 -0 381051267	0.172489573 0.851828047	-0.912767 -2 050604	-0.236619841 1.288501026	-3.331756125 -0 447333553	0.00086			
Total between				-				15.1847909	c,	0.002
Overall	47	-0.782161057	0.105835916	-0.989596	-0.574726474	-7.390317864	1.5E-13			
Mixed effects analysis of extent $(\mathrm{km}^2)$	extent (km²)									
<0.01	21	-1.706183548	0.370467508	-2.432287	-0.980080574	-4.605487687	4.1E-06			
0.01~1	7	-0.582461722	0.408150694	-1.382422	0.217498939	-1.427075172	0.15356			
1~100	4	-0.435016063	0.176635013	-0.781214	-0.0888178	-2.462796342	0.01379			
100~10000	12	-0.811400437	0.162402514	-1.129704	-0.49309736	-4.996230783	5.8E-07			
>10000	3	-0.25454333	0.315088266	-0.872105	0.363018324	-0.807847695	0.41918			
Total within										
Total between								12.1306208	4	0.016
Overall	47	-0.677172027	0.1035127	-0.880053	-0.474290862	-6.541922146	6.1E-11			
Mixed effects analysis of temporal scales	temporal scales									
long-term	1	-2.997706948	0.505159926	-3.987802	-2.007611687	-5.9341741	3E-09			
short-term	46	-1.037141369	0.171330016	-1.372942	-0.701340708	-6.05347149	1.4E-09		-	Fo Lo
I otal between		1 930300647	12025001	1 667407	0 031300405	7 62070763	0 0E 14	4/C88UC.51	_	ZE-04
			1004044010		0010/012/.0	00011	11-17-7			
Mixed effects analysis of species	species									
Acacia saligna	3	-4.443252655	0.772123062	-5.956586	-2.929919261	-5.754591299	8.7E-09			
Acacia longifolia	12	-0.574693192	0.172489573	-0.912767	-0.236619841	-3.331756125	0.00086			
Acacia melanoxylon	-	-7.689849727	1.832127026	-11.28075	-4.09894674	-4.197225201	2.7E-05			
Pinus pinaster	1	-0.730112887	0.540120758	-1.78873	0.328504346	-1.351758613	0.17645			
Different	3	-2.256375532	0.49438226	-3.225347	-1.287404107	-4.564030132	5E-06			
Asparagus asparagoides	Ι	-2.11805426	0.558707299	-3.2131	-1.023008075	-3.790990851	0.00015			
Ailanthus altissima	. 5	-0.298376486	0.138759776	-0.570341	-0.026412323	-2.150309655	0.03153			
Carpobrotus spp.	9 1	-0.933931414	0.158401473	-1.244393	-0.623470231	-5.89597681	3.7E-09			
Uxalis pes-caprae	1	-0.677334604	0.283398874	-1.232786	-0.121883017	-2.390039852	C8910.0			

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e —	-0.891940192 -0.684650429	0.514439811	-1.422876 -1.692934	-0.36100453 0.323633072	-3.2926224 -1.330865953	0.00099 0.18323	65.9736814 11	7E-10
47	-0.732265113	0.076846942	-0.882882	-0.581647875	-9.528877702	0		
Mustralia Lustralia	-1.632729308	0.501253496	-2.615168	-0.650290509	-3.25729261	0.00112		
0 m	-1.47721096 -1765041586	0.190694969	-2.138797	0.041240400 -1 391286315	-1.906/3224/	0.0000		
16	-0.777383315	0.136888616	-1.04568	-0.509086558	-5.678947892	1.4E-08		
Ω.	-0.298376486	0.138759776	-0.570341	-0.026412323	-2.150309655	0.03153		
4	-0.381051267	0.851828047	-2.050604	1.288501026	-0.447333553	0.65463	42.4013174 5	5E-08
47	-0.822482264	0.084567953	-0.988232	-0.656732121	-9.725696683	0		
Mixed effects analysis of invaded country	Ň							
7	-3.409153546	0.710910135	-4.802512	-2.015795286	-4.795477487	1.6E-06		
30	-0.566066113	0.089512527	-0.741507	-0.390624784	-6.323875908	2.6E-10		
- 0	-2.11805426 1 03000253	642101866.0	-3.2131 1 04071	C10200020.1-	-3.1909900851 2 520077720	CIUUU.U		
ĸ	000700400.1-	0.41240000	17040.1-	101040107.0-	01111070-7-	0.0110		
							23.7111967 3	3E-05
47	-0.664560877	0.085791756	-0.83271	-0.496412125	-7.746209056	9.5E-15		
Mixed effects analysis of growth forms								
23	-1.287094996	0.340359632	-1.954188	-0.620002374	-3.781573586	0.00016		
1	-2.11805426	0.558707299	-3.2131	-1.023008075	-3.790990851	0.00015		
7	-0.825590489	0.174204444	-1.167025	-0.484156054	-4.739204538	2.1E-06		
10	-0.786027168	0.221188511	-1.219549	-0.352505653	-3.553652788	0.00038		
4 0	-0.381051267	0.851828047	-2.050604	1.288501026	-0.44/333553	0.65463		
Z	1646410277-	0.010041390	+C70/C.C-	-0.724344971	010070970.0-	0.00088	10.529787 5	0.062
47	-0.968281736	0.120569681	-1.204594	-0.731969504	-8.030889112	8.9E-16		
Mixed effects analysis of habitat types								
15	-2.049044039	0.474893787	-2.979819	-1.118269319	-4.314741726	1.6E-05		
12	-0.574693192	0.172489573	-0.912767	-0.236619841	-3.331756125	0.00086		
18	-0.604948437	0.121655475	-0.843389	-0.366508088	-4.972636354	6.6E-07		
-	-0.684650429	0.514439811	-1.692934	0.323633072	-1.330865953	0.18323		
1	-1.832082371	0.284812292	-2.390304	-1.273860537	-6.432595869	1.3E-10		
							24.3058357 4	7E-05
47	-0.775960769	0.09064049	-0.953613	-0.598308672	-8.560862425	0		



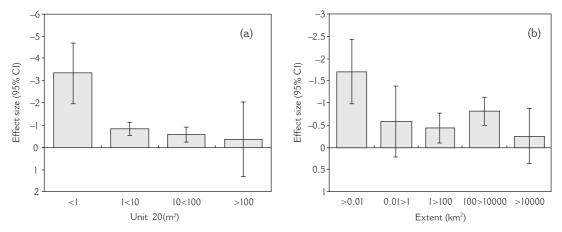
**Figure 2** Meta-analysis of species richness of sites invaded by alien species and their natural reference community in four different Mediterranean-type ecosystems. Mean difference effect size, g, and a mixed (random) effects model (indicated in grey) were used. Case studies 1–47 are arranged from top to bottom (see Table 1)

#### **IV** Discussion

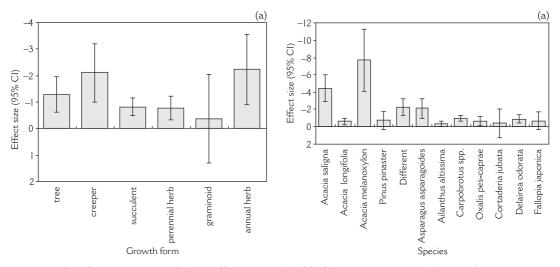
The impact of biological invasions on native ecosystems is the subject of ongoing debate in the literature. Many authors associate invasions with biodiversity declines (Pyšek and Pyšek, 1995; Higgins *et al.*, 1999; Seabloom *et al.*, 2003; French *et al.*, 2008; Hejda *et al.*, 2009). However, some authors argue that the number of naturalized species far exceeds the number of extinctions and that, on balance, introductions over the past few centuries has increased regional biodiversity levels (Rosenzweig, 2001; Davis, 2003; Brown and Sax, 2004; Gurevitch and Padilla, 2004; Houlahan and Findlay, 2004).

The studies reviewed in this paper show that plant invasions are often associated with a significant decline of native plant species richness. This decline varies according to different spatial and temporal scales. The impact of invasive alien species on native species richness is stronger at small spatial scales and decreases with increasing extent of study area and unit size, respectively. The reason for this is that studies at small scales are more likely to detect effects of competition (Huston, 1999) whereas studies conducted over larger areas are more likely to detect the effects of extrinsic factors (mean sitewide biotic or abiotic factors that covary with biodiversity) (Levine and D'Antonio, 1999). Some theories predict that at larger scales increased heterogeneity in resource availability and site conditions may favour the coexistence of native and invasive alien species, provided that they have different functional traits, competitive ability and resource optima (Davies et al., 2005; Smith and Shurin, 2006; Melbourne et al., 2007). Taking this theory further, one could argue that high heterogeneity at large spatial scales promotes diversity of both native and alien species. However, patterns of species diversity at larger scales (ie, regional or subglobal scales) do not necessarily reflect the impact on local biotic interactions (Smith and Shurin, 2006). Our meta-analysis clearly shows that invasions can reduce species richness at small scales.

Another important factor is temporal scale. Investigations in areas with a long invasion history revealed a much stronger impact of invasive alien species on native species

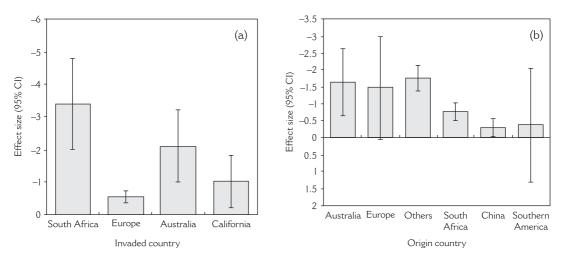


**Figure 3** Effect size (95% Cl) of invasion on species richness for different (a) unit sizes (m<sup>2</sup>) and (b) extents (km<sup>2</sup>) in Mediterranean-type ecosystems. Q-test shows significant different effect sizes (heterogeneity) between groups (a: Q = 15.18, p < 0.01; b: Q = 12.13, p = 0.02)



**Figure 4** Comparison of the effect size (95% Cl) on species richness from species with different (a) growth forms and (b) taxonomical groups in Mediterranean-type ecosystems. Q-test shows no significant different among growth forms (Q = 10.53; p = 0.062), but significant difference of effect sizes (heterogeneity) between species (Q = 65.97; p < 0.001)

richness than studies in recently invaded areas. This finding is supported by a long-term study from permanent plots which demonstrates a decline of plant species richness over a 10-year timeframe of *Lonicera japonica* invasion (Yurkonis and Meiners, 2004). Richardson *et al.* (2007) argued that a timelag between invasions and extinctions could be the reason for the temporary increase of species richness after invasions. Timelags in extinctions could create, and have already created, a large extinction debt which could



**Figure 5** Comparison of the effect size (95% Cl) of invasion on species richness in (a) different invaded countries and (b) the effect size of invasion from species with different countries of origin in Mediterranean-type ecosystems. Q-test shows significant different effect sizes (heterogeneity) between groups (a: Q = 23.7, p < 0.001; b: Q = 42.4, p < 0.001)

be paid in future even with no further introduction of alien plant species (Tilman *et al.*, 1994; Sax and Gaines, 2008). Processes of extinction debt have also been recorded for extinctions related to habitat destruction or fragmentation (Helm *et al.*, 2006). Another interesting aspect is that invasive alien species might not have a direct impact on extinction rates through competitive displacement of established plant species but rather influence colonization rates, thus leading to declines of local diversity (Yurkonis and Meiners, 2004).

Levine *et al.* (2003), in their review of mechanisms underlying the impact of alien plant invasions, posit that it is very difficult to uncover simple rules concerning which invaders or functional groups are most likely to exert large impacts across systems, or which communities will be most susceptible to impacts. Hejda *et al.* (2009) found that a decrease in species richness after alien invasion was largely driven by the identity of the invading species. We could detect patterns relating to the most successful growth forms among invaders and relating to the types of native habitats most affected

by invaders. Shrublands were significantly affected by alien invasion. Trees are the growth form with the highest impact on native species richness, whereas graminoids caused insignificant reductions. These results are consistent with findings from Mason et al. (2009), who found a strong negative effect of woody invaders but little effect of graminoids on shrub species richness. Shrublands in MTEs generally lack tree species and therefore a decline of species richness might be caused by canopy-level changes due to tree invasions. Tree invasion results in higher canopy cover (Rejmánek, 1989), which might lead to species declines through shading effects. Investigations in the South African fynbos and Australian kwongan suggest that even high densities of indigenous overstorey shrubs (Proteaceae species) lead to a decrease in species richness of native plant species (Specht and Specht, 1989; Cowling and Gxaba, 1990; Vlok and Yeaton, 2000). The comparatively low impact of graminoids on native species richness in our meta-analysis was unexpected. Invasive alien grasses are seen as a serious challenge, especially in North America (Seabloom *et al.*, 2003; Callaway and Ridenour, 2004; Moyes *et al.*, 2005; Seastedt and Suding, 2007) but also in other parts of the world (Clarke *et al.*, 2005; Musil *et al.*, 2005). The decline of plant species richness is obviously only one measure of the impact of invasions on invaded ecosystems. The indirect effect of invasive grasses on floristic composition by changing fire regimes is widely recognized as a significant ecological factor (Brooks *et al.*, 2004; Clarke *et al.*, 2005; Rossiter-Rachor *et al.*, 2008).

Old fields showed significant declines in species richness after invasion. Anthropogenically disturbed habitats have been described as habitats with highest frequency and number of alien species (Vilà *et al.*, 2007). Invasions in old fields often hamper successional dynamics and old fields remain in a degraded state once invaded as alien species establish persistent communities that prevent the establishment of native species (Cramer *et al.*, 2008).

Forest habitats were less affected by alien invasions than other habitat types. In general, undisturbed and successionally advanced communities are less invasible than other habitats (Rejmánek, 1989). This finding is supported by studies in Poland (Knight et al., 2008) and Australia (Mason et al., 2009). Introduced Australian Acacia species caused the most severe decline of native species richness, with South Africa being the most affected country. Invasions of Australian Acacia species in fynbos are of particular concern. Australian Acacia species have a huge invasive potential and strong persistence due to enormous loads of long-lived seeds (Richardson and Kluge, 2008). They therefore have radically increased biomass and changed fuel properties in fynbos ecosystems (van Wilgen and Richardson, 1985). Apart from this, Acacia species have massive influences due to nitrogen fixation (Yelenik et al., 2004). These factors in combination have wreaked havoc on fynbos communities.

One could conclude that species introduced from regions with similar climates within MTEs have much stronger impacts, with the native species richness in South Africa declining most severely due to invasion by Australian *Acacia* species. However, this conclusion might be premature as this cohesion could simply be caused by the fact that Australian *Acacia* species were not distributed equally to all the investigated countries.

When investigating declines of native species richness due to alien invasions one has to consider that in many instances invasive alien species have been found to be symptomatic of land-use change (Maskell et al., 2006). In other words, many invaded systems are heavily impacted by habitat loss and disturbance (MacDougall and Turkington, 2005). This led to the reasoning that invasive species might be 'passengers' of degraded ecosystems rather than acting as drivers of degradation. One theory of invader success is that alien species fill unoccupied niches after extinction of native species due to degradation (Shea and Chesson, 2002). Because there might be pre-existing differences prior to invasion, it is difficult to attribute extinctions exclusively to the influence of alien plants.

Indeed most threatened species face more than one threat. It is difficult to disentangle the proximate and ultimate causes of decline or interactions between threats and to evaluate their relative importance (Gurevitch and Padilla, 2004). For example, habitat loss has been identified as the primary cause of extinctions at local and meta community levels in most areas of the world (Davis, 2003). Furthermore, an assessment of threats on biodiversity in the Mediterranean biome showed that both threatened mammals and plants had a negative correlative relationship with the amount of available natural area. with more species threatened when less area remained (Underwood et al., 2009). The question of which factor is most responsible for species declines and which of the factors are drivers or passengers is secondary as ultimately global biodiversity is changing at an

unprecedented rate (Sala *et al.*, 2000) and it is crucial to minimize any impacts.

To understand impacts of alien invasions on native ecosystems it is important to investigate underlying mechanisms. Studies on impacts of invasive species on ecosystem processes concentrate mainly on one mechanism at a time. There are studies which investigate changes of above-ground vegetation due to alien plant invasion (Holmes, 1990; D'Antonio and Mahall, 1991; Blanchard and Holmes, 2008). Other studies concentrate on changes in the soil seed bank (Wearne and Morgan, 2006; Fourie, 2008; Vosse et al., 2008), or changes of soil chemical properties (Witkowski, 1991; Musil, 1993; Yelenik et al., 2004; Lindsay and French, 2005). A range of papers look at competitive interactions between native and invasive plant species (D'Antonio and Mahall, 1991; Sans et al., 2004; Garcia-Serrano et al., 2007; French et al., 2008), while others focus on allelopathic mechanisms (Ridenour and Callaway, 2001; Bais et al., 2003) or a change in soil microorganisms (Allsopp and Holmes, 2001). Assumptions about correlations between changes of vegetation structure and composition and the above-mentioned factors have been made (Holmes, 1990; Musil, 1993). However, our understanding about the interactions between the different mechanisms remains rudimentary. Further research should concentrate on mechanisms underlying alien plant invasions to get a better understanding about which factors are ultimately responsible for a decrease of native species richness.

The meta-analysis approach, although most useful for uncovering the patterns described above, clearly has some limitations that must be considered when evaluating the patterns that have emerged in this study. One problem relates to a potential publication bias in favour of studies that show strong negative impacts on biodiversity: we suggest that studies demonstrating significant impacts are more likely to be published overall than those reporting insignificant impacts. We feel that we have reduced the effect of this potential bias to some extent by including studies from a wide range of journals (ranging from top-tier publications to those with low impact factors), rather than limiting the review to papers in a specific set of journals (high-impact journals are more likely to report dramatic effects, whereas studies with non-significant effects are more likely to be published in journals with lower impact factors). Furthermore, meta-analysis, to a large degree, eliminates bias caused by significant studies with low sample sizes and high sample variance. Nonetheless, some invasive plant species have little or no de-tectable impact (examples of such 'benign invaders' are listed in Richardson et al. (2000: 101), which contribute much weight to the metaanalysis, especially when species richness shows little variation over a large number of samples. Given that, we feel that metaanalysis can improve the robustness of syntheses and should be used in further studies on the mechanisms causing biodiversity decline and biotic homogenization.

Another limitation which has to be considered is the fact that the studies which were included in the meta-analysis all use space for time substitution. The comparison of invaded and uninvaded sites introduces some uncertainty regarding the character of the invaded site prior to invasion.

#### V Conclusions

We have confirmed that in most cases where the effects of plant invasions on native plant diversity have been assessed in Mediterranean-type ecosystems, there are clear negative impacts. It has often been acknowledged that the type and magnitude of impacts depend on the spatial and temporal scale (Sax *et al.*, 2002; Davies *et al.*, 2005). However, our study highlights the importance of the growth form of the invading species, the invaded habitat, as well as the area of investigation. Our study confirmed that invasions indeed cause a marked decline of native plant species richness. Since most of the invasions that were included in our meta-analysis are fairly recent (mostly a few decades) and the results show that the magnitude of impacts increases markedly over time, there is no doubt that declines in species richness is likely to escalate rapidly. These results provide further motivation for urgent action to reduce the extent of alien plant invasions in Mediterranean-type ecosystems.

#### Acknowledgements

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